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(54) Title: HIV ENVELOPE POLYPEPTIDES AND VACCINE

(57) Abstract

Oligonucleotide sequences encoding gp120 polypeptides from breakthrough isolates of vaccine trials using MN-rgp120 and the encoded gp120 polypeptides are provided. Use of the gp120 polypeptides from one or more of the isolates in a subunit vaccine, usually together with MN-rgp120, can provide protection against HIV strains that are sufficiently different from the vaccine strain (e.g.; MN-rgp120) that the vaccine does not confer protection against those strains. Antibodies induced by the polypeptides are also provided.

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## HIV ENVELOPE POLYPEPTIDES AND VACCINE

BACKGROUND OF THE INVENTION5 Field of the Invention

This invention relates to HIV envelope polypeptides and vaccines containing the polypeptides.

Description of the Related Art

10 Acquired immunodeficiency syndrome (AIDS) is caused by a retrovirus identified as the human immunodeficiency virus (HIV). There have been intense efforts to develop a vaccine that induces a protective immune response based on induction of antibodies or 15 cellular responses. Recent efforts have used subunit vaccines where an HIV protein, rather than attenuated or killed virus, is used as the immunogen in the vaccine for safety reasons. Subunit vaccines generally include gp120, the portion of the HIV envelope protein 20 which is on the surface of the virus.

The HIV envelope protein has been extensively described, and the amino acid and nucleic acid sequences encoding HIV envelope from a number of HIV strains are known (Myers, G. et al., 1992. Human 25 Retroviruses and AIDS. A compilation and analysis of nucleic acid and amino acid sequences. Los Alamos National Laboratory, Los Alamos, New Mexico). The HIV envelope protein is a glycoprotein of about 160 kd (gp160) which is anchored in the membrane bilayer at 30 its carboxyl terminal region. The N-terminal segment, gp120, protrudes into the aqueous environment surrounding the virion and the C-terminal segment, gp41, spans the membrane. Via a host-cell mediated process, gp160 is cleaved to form gp120 and the 35 integral membrane protein gp41. As there is no covalent attachment between gp120 and gp41, free gp120 is sometimes released from the surface of virions and

infected cells.

The gp120 molecule consists of a polypeptide core of 60,000 daltons which is extensively modified by N-linked glycosylation to increase the apparent molecular weight of the molecule to 120,000 daltons.

5 The amino acid sequence of gp120 contains five relatively conserved domains interspersed with five hypervariable domains. The positions of the 18 cysteine residues in the gp120 primary sequence, and 10 the positions of 13 of the approximately 24 N-linked glycosylation sites in the gp120 sequence are common to all gp120 sequences. The hypervariable domains contain extensive amino acid substitutions, insertions and deletions. Sequence variations in these domains result 15 in up to 30% overall sequence variability between gp120 molecules from the various viral isolates. Despite this variation, all gp120 sequences preserve the ability of the virus to bind to the viral receptor CD4 and to interact with gp41 to induce fusion of the viral 20 and host cell membranes.

gp120 has been the object of intensive investigation as a vaccine candidate for subunit vaccines, as the viral protein which is most likely to be accessible to immune attack. At present, clinical 25 trials using gp120 MN strain are underway. However, to date no human vaccine trial has been of sufficient size to confirm or refute vaccine efficacy.

The development of candidate HIV-1 vaccines is burdened by the lack of in vivo or in vitro models of 30 HIV-1 infection that accurately approximate the conditions of natural infection in humans. Several candidate HIV-1 vaccines [Berman et al.; *J. Virol.* 7:4464-9 (1992); Haigwood et al.; *J. Virol.* 66:172-82 (1992); Salmon-Ceron et al.; *AIDS Res. and Human 35 Retroviruses* 11:1479-86 (1995)] have been described that elicit broadly cross-reactive antibodies able to

neutralize a variety of diverse HIV-1 isolates *in vitro*. However, the relevance of *in vitro* assays to protective immunity *in vivo* is uncertain. Although several vaccines have provided chimpanzees with protection from challenge by homologous and heterologous strains of HIV-1, protection has not always correlated with *in vitro* neutralization assays carried out in T cell lines, or in lectin- and cytokine-activated peripheral blood mononuclear cells (PBMCs) (Berman et al.; *Nature* 345:622-5 (1990); Bruck et al.; *Vaccine* 12(12):1141-8 (1994); El-Amad et al.; *AIDS* 9:1313-22 (1995); Girard et al.; *J. Virol.* 69:6239-48 (1995); and Fulz et al.; *Science* 256:1687-1690 (1992)]. While successful protection of chimpanzees is encouraging and has historically proved to be a reliable indicator of vaccine efficacy, the conditions of infection in all experimental models of HIV-1 infection differ significantly from natural infection in humans.

Experimental HIV-1 infection *in vivo* and *in vitro* both suffer from the limitation that the *in vitro* amplification of HIV-1, which is required to prepare virus stocks for *in vitro* or *in vivo* infectivity experiments, imposes a genetic selection that results in a spectrum of virus quasi-species that differ from the spectrum of variants present in the clinical specimens used to establish the culture [Kusumi et al.; *J. Virol.* 66:875 (1992); Meyerhans et al.; *Cell* 58:901-10 (1989)]. Because of these uncertainties, and even greater uncertainties related to the amount of virus transmitted, the site and cell type involved in initial replication, and the kinetics of virus dissemination, the ability of currently available *in vitro* or *in vivo* assays to reliably predict vaccine efficacy is questionable.

One of the candidate HIV-1 vaccines that have

entered human clinical trials is recombinant gp120 prepared in Chinese hamster ovary (CHO) cells from the MN strain of HIV-1 (MN-rgp120) (Berman et al.; *J. Virol.* 7:4464-9 (1992)). To date, approximately 499 5 adults have participated in Phase 1 and 2 immunogenicity and safety trials of this vaccine. The data collected thus far suggest that MN-rgp120 is safe, immunogenic, and elicits high titers of neutralizing antibodies in greater than 95% of individuals immunized 10 according to a 0, 1, and 6 month immunization schedule [Belshe et al.; *JAMA* 272(6):475-80 (1994); McElrath; *Seminars in Cancer Biol.* 6:1-11 (1995)]. However, 15 during the course of these trials, nine vaccinees who received MN-rgp120 have become infected with HIV-1 through high risk behavior. Small trials, such as these, in populations with low rates of infection and minimally sized placebo control groups do not have sufficient statistical power to confirm or refute 20 vaccine efficacy.

However, effective vaccines based on gp120 or another HIV protein for protection against additional strains of HIV are still being sought to prevent the spread of this disease.

25 Description of the Background Art

Recombinant subunit vaccines are described in Berman et al., PCT/US91/02250 (published as number WO91/15238 on 17 October 1991). See also, e.g., Hu et al., *Nature* 328:721-724 (1987) (vaccinia virus- 30 HIV envelope recombinant vaccine); Arthur et al., *J. Virol.* 63(12): 5046-5053 (1989) (purified gp120); and Berman et al., *Proc. Natl. Acad. Sci. USA* 85:5200-5204 (1988) (recombinant envelope glycoprotein gp120).

35 Numerous sequences for gp120 are known. The sequence of gp120 from the IIIB substrain of HIV-1

referred to herein is that determined by Muesing et al., "Nucleic acid structure and expression of the human AIDS/lymphadenopathy retrovirus, *Nature* 313:450-458 (1985). The sequences of gp120 from the 5 NY-5, JRCSF, 26, Z321, and HXB2 strains of HIV-1 are listed by Myers et al., "Human Retroviruses and AIDS; A compilation and analysis of nucleic acid and amino acid sequences," Los Alamos National Laboratory, Los Alamos, New Mexico (1992). The sequence of the Thai isolate 10 A244 is provided by McCutchan et al., "Genetic Variants of HIV-1 in Thailand," *AIDS Res. and Human Retroviruses* 8:1887-1895 (1992). The MN<sub>194</sub> clone is described by Gurgo et al., "Envelope sequences of two new United States HIV-1 isolates," *Virology* 164: 531-536 (1988). As 15 used herein, MN, MN-rgp120, the MN clone or isolate refers to MN<sub>GNE</sub>. The MN<sub>GNE</sub> amino acid sequence is Sequence ID No. 29.

Each of the above-described references is 20 incorporated herein by reference in its entirety.

#### Summary of the Invention

Oligonucleotide sequences encoding gp120 polypeptides from breakthrough isolates of vaccine trials using MN-rgp120 and the encoded gp120 25 polypeptides are provided. Use of the gp120 polypeptides from one or more of the isolates in a subunit vaccine, usually together with MN-rgp120, can provide protection against HIV strains that are sufficiently different from the vaccine strain (e.g.; 30 MN-rgp120) that the vaccine does not confer protection against those strains. Antibodies induced by the polypeptides are also provided.

#### Brief Description of the Drawings

35 Figure 1 illustrates the kinetics of antibody response to MN-rgp120 in vaccinees infected with HIV-1.

Sera were collected at the time points indicated and assayed for antibodies reactive with MN-rgp120 (open circles) or a synthetic peptide derived from the V3 domain of MN-rgp120 (closed circles). Arrows indicate 5 dates of injection. Plus sign indicates the first time HIV-1 infection was detected. Shaded area indicates data collected after HIV-1 infection. Data from vaccinee C6 is shown in panel A; C8 in panel B; C7, panel C; C11, panel D; C10, panel E; C17, panel F; and 10 C15, panel G.

Figure 2 illustrates the kinetics of CD4 blocking antibody response in vaccinees infected with HIV-1. Sera were collected at the time points indicated and assayed for antibodies able to block the binding of 15 [<sup>125</sup>I]-labeled MN-rgp120 to cell surface CD4. Arrows indicate dates of injection. Plus sign indicates the first time HIV-1 infection was detected. Shaded area indicates data collected after HIV-1 infection. Data from vaccinee C6 is shown in panel A; C8 in panel B; 20 C7, panel C; C11, panel D; C10, panel E; C17, panel F; and C15, panel G.

Figure 3 illustrated predicted amino acid sequences of envelope glycoproteins (gp120) from breakthrough viruses. Proviral DNA sequences were 25 amplified by PCR from PBMCs and cloned into the PRK5 expression plasmid. Two clones from each infected vaccinee were sequenced from double stranded plasmid DNA. Sequence numbering is with reference to the initiator methionine residue of gp120. For the purpose 30 of comparison, the sequences shown begin at amino acid 12 of the mature, fully processed, envelope glycoproteins (corresponding to position 41 of the gp120 open reading frame). Shaded areas indicate sequences at neutralizing epitopes, dark boxes indicate 35 polymorphisms thought to be important for the binding of virus neutralizing MAbs reactive with MN-rgp120.

conserved (C) regions and variable (V) regions are indicated above the sequences. Boxes indicate sequence homologies and polymorphisms.

Figure 4 illustrates immunoprecipitation of recombinant gp120 prepared from breakthrough viruses.

Recombinant gp120s from the seven breakthrough viruses were prepared by transient transfection of 293s cells. Cells were metabolically labeled with  $^{35}$ S methionine and growth conditioned cell culture supernatants were immunoprecipitated with polyclonal antisera to MN-rgp120. Immunoprecipitates were resolved by SDS-PAGE and visualized by autoradiography. C8 lanes a and b correspond to clones C8.3 and C8.6; C6 lanes a and b correspond to clones C6.1 and C6.5; C7 lanes a and b correspond to clones C7.2 and C7.10; C17 lanes a and b correspond to clones C17.1 and C17.3; C11 lanes a and b correspond to clones C11.5 and C11.7; C10 lanes a and b correspond to clones C10.5 and C10.7; C15 lanes a and b correspond to clones C15.2 and C15.3.

Figure 5 illustrates binding of monoclonal antibodies to recombinant gp120 from breakthrough viruses. Growth-conditioned cell culture supernatants were collected from 293s cells transiently transfected with plasmids directing the expression of breakthrough virus envelope glycoproteins. The relative rgp120 concentrations were determined by ELISA using MAb 5B6 specific for the HSV-1 glycoprotein D flag epitope at the amino terminus of all of the rgp120 variants described herein. The resulting rgp120 preparations were captured onto wells of microtiter plates coated with a polyclonal antibody specific for a conserved sequence in the C-terminus of gp120. The binding of virus neutralizing monoclonal antibodies reactive with gp120 was determined by ELISA. A, binding by MAb (5B6) specific for the HSV-1 glycoprotein D flag epitope; B, binding by MAb (1034) against the V3 domain of

MN-rgp120; C binding by MAb (50.1) raised against a synthetic peptide corresponding to the V3 domain of MN-rgp120; D, binding by a human MAb (15e) known to block the binding of gp120 to CD4.

5 Figure 6 depicts the mature envelope glycoprotein (gp120) from the MN clone of the MN strain of HIV-1 (SEQ. ID. NO. 29). Hypervariable domains are indicated in bold, and the V and C regions are indicated (according to Modrow et al., *J. Virology* 61(2):570  
10 15 (1987). Potential glycosylation sites are marked with a (\*).

#### Detailed Description of the Invention

The present invention provides gp120 polypeptides from breakthrough isolates of HIV vaccine trials. 15 Novel oligonucleotide sequences encoding gp120 from breakthrough isolates which can be used to express gp120 are also provided. Use of gp120 polypeptides from one or more of the isolates in a subunit vaccine, 20 usually together with MN-rgp120, can provide protection against HIV strains that are sufficiently different from the vaccine strain (e.g.; MN-rgp120) that the vaccine does not confer protection against those strains.

25 In one embodiment, the vaccine is based on the use of the MN-rgp120 polypeptide (Sequence ID No. 29) and gp120 polypeptides from MN-like viruses that include neutralizing epitopes that are not present in the initial vaccine strain, and are sufficiently different 30 from those of the vaccine strain, to have been able to cause HIV-1 infections in MN-rgp120 vaccinated individuals (i.e., to result in breakthrough infections). Use of the initial vaccine strain empirically determines the viruses present in the 35 population that contain additional neutralizing epitopes sufficiently different from those of the

vaccine strain to escape protection induced by the vaccine strain. Use of an initial representative gp120 polypeptide in a vaccine acts as a sieve so that viruses that are not effectively protected against by the vaccine strain breakthrough the vaccine, empirically resulting in determination of additional strains in a given geographic region that are not protected against by the initial vaccine strain. Use of gp120 from those breakthrough isolates complements the vaccine isolate by providing additional neutralizing epitopes not present in the initial vaccine strain, therefore creating a more complete vaccine that confers protection against multiple different virus strains in the region.

Prior HIV-1 vaccine strategies were based on selection of appropriate candidate vaccine polypeptides based on homology alignment studies. However, since some of the neutralizing epitopes are conformation-dependent and the location of all of these epitopes is not known, this approach necessarily cannot determine all of the neutralizing epitopes that should be included in a vaccine for a particular region. In contrast, the present approach uses a selected representative strain and empirically determines strains that are sufficiently different and therefore breakthrough the barrier of protection provided by the initial vaccination program. Those strains can be included in the vaccine to confer more complete protection from HIV strains in the region. In addition, those strains can be used alone to confer protection against the breakthrough virus.

In another embodiment, the invention comprises a vaccine containing a first HIV gp120 polypeptide sequence and a breakthrough isolate HIV gp120 polypeptide sequence from a vaccinee vaccinated with a vaccine including the first HIV gp120 polypeptide.

sequence, the HIV gp120 polypeptide sequences being in a suitable carrier. Fragments of one or both HIV gp120 polypeptide sequences can be substituted for one or both of the corresponding HIV gp120 polypeptide sequences.

Preferably, the first gp120 polypeptide sequence contains neutralizing epitopes found in one or more gp120 polypeptides present in isolates from the geographical region where the initial vaccine (i.e., 10 the vaccine that gives rise to the breakthrough isolate) is administered. More preferably, the first gp120 polypeptide sequence contains at least one of the more common neutralizing epitopes for the region, and most preferably the first gp120 polypeptide sequence 15 contains at least one of the three most common neutralizing epitopes.

gp120 polypeptide sequences suitable for use as the first gp120 polypeptide sequence include gp120 MN, the Thai isolate A244 sequence (hereinafter "gp120 20 A244"), gp120 MN-GNE6 (Sequence ID No. 31; also known in the art as "gp120 GNE6"), and gp120 MN-GNE8 (Sequence ID No. 33; also known in the art as "gp120 GNE8"), and the like. gp120 MN, gp120 MN-GNE6, and gp120 MN-GNE8 are especially preferred for use as the 25 first gp120 polypeptide sequence in initial vaccines for North America. gp120 A244 is especially preferred for use as the first gp120 polypeptide sequence in initial vaccines for Thailand.

In a variation of this embodiment, the vaccine 30 includes two different (i.e., first and second) gp120 polypeptide sequences, or fragments thereof, in combination with a breakthrough isolate HIV gp120 polypeptide sequence. The latter can be from a vaccinee vaccinated with either or both of the first 35 and second HIV gp120 polypeptide sequences.

Exemplary vaccines include those containing

combinations of gp120 MN, gp120 A244, gp120 MN-GNE6 (Sequence ID No. 31), and gp120 MN-GNE8 (Sequence ID No. 33). Combinations of gp120 MN and gp120 A244 or gp120 MN-GNE8 (Sequence ID No. 33) with a breakthrough isolate HIV gp120 polypeptide sequence are especially preferred.

5 In vaccines containing gp120 MN, the breakthrough isolate HIV gp120 polypeptide sequence can be an HIV gp120 polypeptide sequence selected from the group consisting of Sequence ID Nos. 2, 4, 6, 8, 10, 12, 14, 10 16, 18, 20, 22, 24, 26, and 28, and fragments thereof.

15 The term "subunit vaccine" is used herein, as in the art, to refer to a viral vaccine that does not contain virus, but rather contains one or more viral proteins or fragments of viral proteins. As used herein, the term "multivalent", means that the vaccine contains gp120 from at least two HIV isolates having different amino acid sequences.

20 The term "breakthrough isolate" or "breakthrough virus" is used herein, as in the art, to refer to a virus isolated from a vaccinee.

25 The terms "amino acid sequence", "polypeptide sequence", and "polypeptide" are used interchangeably herein as in the art, as are the terms "nucleic acid sequence", "nucleotide sequence", and "oligonucleotide".

#### Polypeptides from Breakthrough Isolates

30 The gp120 polypeptides of this invention correspond to the amino acid sequences of seven breakthrough isolates which are illustrated below in Table 1. A polypeptide of this invention includes an HIV gp120 amino acid sequence illustrated in Table 1 (Sequence ID Nos. 1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 35 21, 23, 25, and 27) and fragments thereof. The polypeptides of this invention can include fused

sequences from two or more HIV gp120 or gp160 amino acid sequences.

The polypeptide can also be joined to another viral protein, such as a flag epitope amino acid sequence. The term "flag epitope" is used herein, as in the art, to denote an amino acid sequence that includes an epitope recognized by a monoclonal antibody. Flag epitopes facilitate using single monoclonal antibody affinity purification of a plurality of different recombinant proteins, each having the flag epitope recognized by the monoclonal antibody. Numerous amino acid sequences can function as flag epitopes. The N-terminal sequences of Herpes Simplex Virus Type 1 (HSV-1) glycoprotein D (gD-1) is conveniently used as the flag epitope and its use is described in detail in the examples. The flag epitope is conveniently fused to the N terminus of the HIV gp120 polypeptide sequence. Alternatively, however, monoclonal antibodies that recognize neutralizing epitopes in the rgp120 sequences can be used to affinity purify the amino acid sequences, and a flag epitope can be omitted.

In addition, various signal sequences can be joined to a polypeptide of this invention. Although rgp120 is secreted to some extent in HIV cultures, the amount of the envelope glycoprotein released from (secreted by) the host cells varies widely from strain to strain. Various signal sequences can be introduced into the polypeptide by joining a nucleotide sequence encoding the signal sequence to the nucleotide sequence encoding the rgp120 to facilitate secretion of rgp120 from the cells. For example, Chiron HIV gp120 polypeptides include a signal sequence from tissue plasminogen activator (TPA) that provides good secretion of rgp120. Additional signal sequences are well known and include the N-terminal domain of murine

leukemia virus surface protein gp70 described by Kayman et al., J. Virol. 68:400-410 (1984).

Table 1 illustrates the nucleotide and deduced amino acid sequences for two clones of each the seven breakthrough isolates of this invention. The clones are: C6.1; C6.5; C8.3; C8.6; C15.2; C15.3; C7.2; C7.10; C11.5; C11.7; C10.5; C10.7; C17.1; and C17.3. These sequences are SEQ. ID. NOS. 1-28, the first sequence number for each clone being the nucleotide sequence and the second being the amino acid sequence. The amino acid sequence for MN and the nucleotide and deduced amino acid sequences for MN-GNE6 and MN-GNE8 are illustrated in the sequence listing hereinafter. In the listing for MN-GNE6, a stop codon appears at amino acid residue position 51. This stop codon can be replaced with a codon encoding the corresponding amino acid from MN or MN-GNE8 or another isolate.

TABLE 1

CLONE C6.1

5 GGG GTC CCT GTG TGG AAG GAA GCA ACC ACC ACT CTA 36  
 Gly Val Pro Val Trp Lys Glu Ala Thr Thr Thr Leu  
 1 5 10  
 TTT TGT GCA TCA GAT GCT AAA GCA TAT GAC ACA GAC CTG 75  
 Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu Val  
 15 20 25  
 10 CAT AAT GTT TGG GCC ACA CAT GCT TGT GTA CCC ACA GAC 114  
 His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp  
 30 35  
 15 CCA AAC CCA CAA GAA ATG GCA TTG GAA AAT GTG ACA GAA 153  
 Pro Asn Pro Gln Glu Met Val Leu Glu Asn Val Thr Glu  
 40 45 50  
 20 GAT TTT AAC ATG TGG AAA AAT GAC ATG GCA GAA CAG ATG 192  
 Asp Phe Asn Met Trp Lys Asn Asp Met Val Glu Gln Met  
 55 60  
 25 CAT GAG GAT ATA ATC AGT TTA TGG GAT CAA AGC CTA AAA 231  
 His Glu Asp Ile Ile Ser Leu Trp Asp Gln Ser Leu Lys  
 65 70 75  
 30 CCA TGT GTA AAA TTA ACC CCA CTC TGT ATT ACT TTA AAT 270  
 Pro Cys Val Lys Leu Thr Pro Leu Cys Ile Thr Leu Asn  
 80 85 90  
 35 TGC ACC AAT TGG AAG AAG AAT GAT ACT AAA ACT AAT AGT 309  
 Cys Thr Asn Trp Lys Lys Asn Asp Thr Lys Thr Asn Ser  
 95 100  
 40 AGT AGT ACT ACA ACT AAT AGT AGT GCT ACA GCA GCT AAT 348  
 Ser Ser Thr Thr Thr Asn Asn Ser Ser Ala Thr Ala Asn  
 105 110 115  
 45 AGT AGT AGT ACT ACA ACT AAT AGT AGT TGG GCA GAG ATA 387  
 Ser Ser Ser Thr Thr Thr Asn Ser Ser Trp Gly Glu Ile  
 120 125  
 50 AAG GAG GGA GAA ATA AAG AAC TGC TCT TTC AAT ATC ACC 426  
 Lys Glu Gly Glu Ile Lys Asn Cys Ser Phe Asn Ile Thr  
 130 135 140  
 55 ACA ACC ATA AGA GAC AAG GTG AAG AAA GAA TAT GCA CTT 465  
 Thr Ser Ile Arg Asp Lys Val Lys Lys Glu Tyr Ala Leu  
 145 150 155  
 60 TTT TAT AGC CTT GAT GTA GTA CCA ATA GAA AAT GAT AAT 504  
 Phe Tyr Ser Leu Asp Val Val Pro Ile Glu Asn Asp Asn  
 160 165  
 65 ACT AGC TAT AGG TTG AGA AGT TGT AAC ACC TCA GTC ATT 543  
 Thr Ser Tyr Arg Leu Arg Ser Cys Asn Thr Ser Val Ile  
 170 175 180  
 70 ACA CAA GCC TGT CCA AAG GTA ACT TTT GAG CCA ATT CCC 582  
 Thr Gln Ala Cys Pro Lys Val Thr Phe Glu Pro Ile Pro  
 185 190  
 75 ATA CAT TAT TGT ACC CCG GCT GGT TTT GCG ATT CTG AAG 621  
 Ile His Tyr Cys Thr Pro Ala Gly Phe Ala Ile Leu Lys  
 195 200 205  
 80 TGT AGA GAT AAA AAG TTC AAT GGA ACA GGA CCA TGC AAA 660  
 Cys Arg Asp Lys Lys Phe Asn Gly Thr Gly Pro Cys Lys  
 210 215 220  
 85 AAT GTT AGC ACA GTA CAA TGT GCA CAT GGA ATT AAG CCA 699  
 Asn Val Ser Thr Val Gln Cys Ala His Gly Ile Lys Pro  
 225 230  
 90 GTA GTG TCA ACT CAA CTG CTG TTA AAT GGC AGC CTA GCA 738  
 Val Val Ser Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala  
 235 240 245  
 95 GAA GAA GAG GTA ATA ATT AGA TCT GCC AAT TTC TCA AAC 777  
 Glu Glu Glu Val Ile Ile Arg Ser Ala Asn Phe Ser Asn  
 250 255

AAT GCT AAA ATC ATA ATA GCA CAG TTG AGG GAA CCT GTA 816  
 Asn Ala Lys Ile Ile Ile Val Gln Leu Arg Glu Pro Val  
 260 265 270  
 GAA ATT AAT TGT ACA AGA CCC AGC AAC AAT ACA ATA AAA 855  
 5 Glu Ile Asn Cys Thr Arg Pro Ser Asn Asn Thr Ile Lys  
 275 280 285  
 GGT ATA CAC ATA GGA CCA GGG AGA GCA TTT TAT GCA ACA 894  
 Gly Ile His Ile Gly Pro Gly Arg Ala Phe Tyr Ala Thr  
 290 295  
 10 GGA GAC ATA CGA GGA GAT ATA AGA CAA GCA CAT TGT AAC 933  
 Gly Asp Ile Arg Gly Asp Ile Arg Gln Ala His Cys Asn  
 300 305 310  
 ATT AGT GGA GCA AAA TGG AAT AAC ACT TTA AAG AAG GTA 972  
 Ile Ser Gly Ala Lys Trp Asn Asn Thr Leu Lys Lys Val  
 15 315 320  
 GTT AAA AAA TTA AAA GAA CAA TTT CCA AAT AAA ACA ATA 1011  
 Val Lys Lys Leu Lys Glu Gln Phe Pro Asn Lys Thr Ile  
 325 330 335  
 20 GTC TTT AAC CAT TCC TCA GGA CGG GAC CCA GAA ATT GTA 1050  
 Val Phe Asn His Ser Ser Gly Gly Asp Pro Glu Ile Val  
 340 345 350  
 ATG CAC AGT TTT AAT TGT CAA GGG GAA TTT TTC TAC TGT 1089  
 Met His Ser Phe Asn Cys Gln Gly Glu Phe Phe Tyr Cys  
 355 360  
 25 AAT ACA ACA AAG CTG TTT AAT AGT ACT TGG AAT GAT ACT 1128  
 Asn Thr Thr Lys Leu Phe Asn Ser Thr Trp Asn Asp Thr  
 365 370 375  
 ACA GAG TCA AAT AAC AAT GAT AGT ACT ATT ACA CTC CCA 1167  
 Thr Glu Ser Asn Asn Asn Asp Ser Thr Ile Thr Leu Pro  
 380 385  
 30 TGC AGA ATA AAA CAA ATT ATA AAC ATG TGG CAG GAA ATA 1206  
 Cys Arg Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Ile  
 390 395 400  
 35 GGA AAA GCA ATG TAT GCC CCT CCC ACC AGA GGA GAA ATT 1245  
 Gly Lys Ala Met Tyr Ala Pro Pro Thr Arg Gly Glu Ile  
 405 410 415  
 AAA TGT TCA TCA AAT ATT ACA GGA CTA CTG TTA ATA AGA 1284.  
 Lys Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Ile Arg  
 420 425  
 40 GAT GGT GGT ATT AAC ACT AGC GAT GCC ACC GAG ACC TTC 1323  
 Asp Gly Gly Ile Asn Thr Ser Asp Ala Thr Glu Thr Phe  
 430 435 440  
 AGA CCG GGA GGA GGA GAT ATG AGG GAC AAT TGG AGA AGT 1362  
 Arg Pro Gly Gly Asp Met Arg Asp Asn Trp Arg Ser  
 445 450  
 45 GAA TTA TAT AAA TAT AAA GTA GTG AAA ATT GAG CCA TTA 1401  
 Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile Glu Pro Leu  
 455 460 465  
 GGA GTA GCA CCC ACC AAG GCA AAG AGA AGA GTG GTG CAG 1440  
 50 Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln  
 470 475 480  
 AGA GAA AAA AGA GCA GTA ACA CTA GGA GCT ATG TTC CTT 1479  
 Arg Glu Lys Arg Ala Val Thr Leu Gly Ala Met Phe Leu  
 485 490  
 55 GGG TTC TTA GGA GCA TAA AGC TTC 1503  
 Gly Phe Leu Gly Ala Xaa Ser Phe  
 495 500 501

CLONE C6.5

60 GGG GTA CCT CTA TGG AAA GAA GCA ACC ACC ACT CTA 36  
 Gly Val Pro Val Trp Lys Glu Ala Thr Thr Leu  
 1 5 10  
 TTT TGT GCA TCA GAT GCT AAA GCA TAT GAC ACA GAG GTG 75  
 Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu Val  
 65 15 20 25

CAT	AAT	GTT	TGG	GCC	ACA	CAT	GCT	TGT	GTA	CCC	ACA	GAC	114
His	Asn	Val	Trp	Ala	Thr	His	Ala	Cys	Val	Pro	Thr	Asp	
						30					35		
5	CCA	AAC	CCA	CAA	GAA	ATG	GTA	TTG	GAA	AAT	GTG	ACA	153
	Pro	Asn	Pro	Gln	Glu	Met	Val	Leu	Glu	Asn	Val	Thr	
						40					45		50
	GAT	TTT	AAC	ATG	TGG	AAA	AAT	GAC	ATG	GTA	GAA	CAG	192
	Asp	Phe	Asn	Met	Trp	Lys	Asn	Asp	Met	Val	Glu	Gln	
10	CAT	GAG	ANT	ATA	ATC	AGT	TTA	TGG	GAT	CAA	AGC	CTA	AAA
	His	Glu	Xaa	Ile	Ile	Ser	Leu	Trp	Asp	Gln	Ser	Leu	Lys
						65				70		75	
15	CCA	TGT	GTA	AAA	TTA	ACC	CCA	CTC	TGT	ATT	ACT	TTA	AAT
	Pro	Cys	Val	Lys	Leu	Thr	Pro	Leu	Cys	Ile	Thr	Leu	Asn
						80				85		90	
	TGC	ACC	AAT	TGG	AAG	GAG	AAT	GAT	ACT	AAA	ACT	AAT	AGT
	Cys	Thr	Asn	Trp	Lys	Glu	Asn	Asp	Thr	Lys	Thr	Asn	Ser
						95					100		
20	AGT	AGT	ACT	ACA	ACT	AAT	AAT	AGT	AGT	GCT	ACA	GCT	AAT
	Ser	Ser	Thr	Thr	Thr	Asn	Asn	Ser	Ser	Ala	Thr	Ala	Asn
						105				110		115	
	AGT	AGT	AGT	ACT	ACA	ACT	AAT	AGT	AGT	TGG	GGA	GAG	ATA
	Ser	Ser	Ser	Thr	Thr	Thr	Asn	Ser	Ser	Trp	Gly	Glu	Ile
						120				125			
25	AAG	GAG	GGA	GAA	ATA	AAG	AAC	TGC	TCT	TTC	AAT	ATC	ACC
	Lys	Glu	Gly	Glu	Ile	Lys	Asn	Cys	Ser	Phe	Asn	Ile	Thr
						130			135			140	
	ACA	GGC	ATA	AGA	GAC	AAG	GTG	AAG	AAA	GAA	TAT	GCA	CTT
	Thr	Gly	Ile	Arg	Asp	Lys	Val	Lys	Lys	Glu	Tyr	Ala	Leu
30						145				150			155
	TTT	TAT	AGC	CTT	GAT	GTA	GTA	CCA	ATA	GAA	AAT	GAT	AAT
	Phe	Tyr	Ser	Leu	Asp	Val	Val	Pro	Ile	Glu	Asn	Asp	Asn
						160					165		
35	ACT	AGC	TAT	AGG	TTG	AGA	AGT	TGT	AAC	ACC	TCA	GTC	ATT
	Thr	Ser	Tyr	Arg	Leu	Arg	Ser	Cys	Asn	Thr	Ser	Val	Ile
						170			175			180	
	ACA	CAA	GCC	TGT	CCA	AAG	GTA	ACT	TTT	GAG	CCA	ATT	CCC
	Thr	Gln	Ala	Cys	Pro	Lys	Val	Thr	Phe	Glu	Pro	Ile	Pro
						185				190			
40	ATA	CAT	TAT	TGT	ACC	CCG	GCT	GGT	TTT	GCG	ATT	CTG	AAG
	Ile	His	Tyr	Cys	Thr	Pro	Ala	Gly	Phe	Ala	Ile	Leu	Lys
						195			200			205	
	TGT	AAA	GAT	AAA	AAG	TTC	AAT	GGA	ACA	GGA	CCA	TGC	AAA
	Cys	Lys	Asp	Lys	Phe	Asn	Gly	Thr	Gly	Pro	Cys	Lys	
45						210				215			220
	AAT	GTT	AGC	ACA	GTA	CAA	TGT	ACA	CAT	GGA	ATT	AAG	CCA
	Asn	Val	Ser	Thr	VaI	Gln	Cys	Thr	His	Gly	Ile	Lys	Pro
						225				230			
50	GTA	GTG	TCA	ACT	CAA	CTG	CTG	TTA	AAT	GGC	AGC	CTA	GCA
	Val	Val	Ser	Thr	Gln	Leu	Leu	Leu	Asn	Gly	Ser	Leu	Ala
						235			240			245	
	GAA	GAA	GAG	GTA	ATA	ATT	AGA	TCT	GCC	AAT	TTC	TCA	AAC
	Glu	Glu	Glu	Val	Ile	Ile	Arg	Ser	Ala	Asn	Phe	Ser	Asn
						250				255			
55	AAT	GCT	AAA	ATC	ATA	ATA	GTA	CAG	TTG	AAG	GAA	CCT	GTA
	Asn	Ala	Lys	Ile	Ile	Ile	Val	Gln	Leu	Lys	Glu	Pro	Val
						260			265			27	
	GAA	ATT	AAT	TGT	ACA	AGA	CCC	AGC	AAC	AAT	ACA	ATA	AAA
	Glu	Ile	Asn	Cys	Thr	Arg	Pro	Ser	Asn	Asn	Thr	Ile	Lys
60						275				280			285
	GGT	ATA	CAC	ATA	GCA	CCA	GGG	AGA	GCA	TTT	TAT	GCA	ACA
	Gly	Ile	His	Ile	Gly	Pro	Gly	Arg	Ala	Phe	Tyr	Ala	Thr
						290				295			

GCA GAC ATA CGA GGA GAT ATA AGA CAA GCA CAT TGT AAC 933  
 Gly Asp Ile Arg Gly Asp Ile Arg Gln Ala His Cys Asn  
 300 305 310  
 ATT AGT GGA GCA AAA TGG AAT AAC ACT TTA AAG AAG GTA 972  
 5 Ile Ser Gly Ala Lys Trp Asn Asn Thr Leu Lys Lys Val  
 315 320  
 GTT ATA AAA TTA AAA GAA CAA TTT CCA AAT AAA ACA ATA 1011  
 Val Ile Lys Leu Lys Glu Gln Phe Pro Asn Lys Thr Ile  
 325 330 335  
 10 GTC TTT AAC CAT TCC TCA GGA GGG GAC CCA GAA ATT GTA 1050  
 Val Phe Asn His Ser Ser Gly Gly Asp Pro Glu Ile Val  
 340 345 350  
 ATG CAC AGT TTT AAT TGT CAA GGG GAA TTT TTC TAC TGT 1089  
 Met His Ser Phe Asn Cys Gln Gly Glu Phe Phe Tyr Cys  
 15 355 360  
 AAT ACA ACG AAG CTG TTT AAT AGT ACT TGG AAT GAT ACT 1128  
 Asn Thr Thr Lys Leu Phe Asn Ser Thr Trp Asn Asp Thr  
 365 370 375  
 ACA GAG TCA AAT AAC AAT GAT AGT ACT ATT ACA CTC CCA 1167  
 20 Thr Glu Ser Asn Asn Asn Asp Ser Thr Ile Thr Leu Pro  
 380 385  
 TGG AGA ATA AAA CAA ATT ATA AAC ATG TGG CAG GAA GTA 1206  
 Cys Arg Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Val  
 390 395 400  
 25 GGA AAA GCA ATG TAT GCC CCT CCC ATC AGA GGA GAA ATT 1245  
 Gly Lys Ala Met Tyr Ala Pro Pro Ile Arg Gly Glu Ile  
 405 410 415  
 AAA TGT TCA TCA AAT ATT ACA GGA CTA CTG TTA ACA AGA 1284  
 Lys Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg  
 30 420 425  
 GAT GGT GGT ATT AAC ACT AGC GAT GCC ACC GAG ACC TTC 1323  
 Asp Gly Gly Ile Asn Thr Ser Asp Ala Thr Glu Thr Phe  
 430 435 440  
 AGA CCG GGA GGA GGA GAT ATG AGG GAC AAT TGG AGA ACT 1362  
 35 Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp Arg Ser  
 445 450  
 GAA TTA TAT AAA TAT AAA GTA GTG AAA ATT GAG CCA TTA 1401  
 Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile Glu Pro Leu  
 455 460 465  
 40 GGA GTA GCA CCC ACC AAG GCA AAG AGA AGA GTG GTG CAG 1440  
 Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln  
 470 475 480  
 AGA GAA AAA AGA GCA GTA ACA CTA CGA GCT ATG TTC CTT 1479  
 Arg Glu Lys Arg Ala Val Thr Leu Gly Ala Met Phe Leu  
 485 490 495  
 45 GGG TTC TTG GGA GCA TAA AGC TTC 1503  
 Gly Phe Leu Gly Ala Xaa Ser Phe  
 500 501  
 50 CLONE C8.3  
 G CTA CCT GTA TGG AAA GAA GCA ACC ACC ACT CTA TTT 37  
 Val Pro Val Trp Lys Glu Ala Thr Thr Leu Phe  
 1 5 10  
 TGT GCA TCA GAT GCT AAA GCA TAT GAT ACA GAG GTA CAT 76  
 55 Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu Val His  
 15 20 25  
 AAT GTT TGG GCT ACA CAT GCC TGT GTA CCC ACA GAC CCC 115  
 Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro  
 30 35  
 60 AAC CCA CAA GAA GTA GTA TTG GAA AAT GTA ACA GAA AAT 154  
 Asn Pro Gln Glu Val Val Leu Glu Asn Val Thr Glu Asn  
 40 45 50  
 TTT AAC ATG TGG AAA AAT AAC ATG GTA GAA CAG ATG CAT 193  
 Phe Asn Met Trp Lys Asn Asn Met Val Glu Gln Met His  
 65 55 60

GAG GAT ATA ATC AGT TTA TGG GAT CAA AGT CTA AAG CCA 232  
 Glu Asp Ile Ile Ser Leu Trp Asp Gln Ser Leu Lys Pro  
 65 70 75  
 TGT GTA AAA TTA ACC CCA CTC TGT GTT ACT TTA AAT TGC 271  
 5 Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu Asn Cys  
 80 85 90  
 ACT AAT TTG GAG AAT GCT AAT ACC GAG AAT GCT AAT 310  
 Thr Asn Leu Glu Asn Ala Asn Asn Thr Glu Asn Ala Asn  
 95 100  
 10 AAT ACC AAT AAT TAT ACC TTG GGG ATG GAG AGA GGT GAA 349  
 Asn Thr Asn Asn Tyr Thr Leu Gly Met Glu Arg Gly Glu  
 105 110 115  
 ATA AAA AAC TGC TCT TTC AAT ATC ACC ACA AGC TTA AGA 388  
 Ile Lys Asn Cys Ser Phe Asn Ile Thr Thr Ser Leu Arg  
 15 120 125  
 GAT AAG GTG AAA AAA GAA TAT GCA TTG TTT TAT AAA CTT 427  
 Asp Lys Val Lys Lys Glu Tyr Ala Leu Phe Tyr Lys Leu  
 130 135 140  
 GAT GTA GTA CAA ATA GAT AAT AGT ACC AAC TAT AGG CTG 466.  
 20 Asp Val Val Gln Ile Asp Asn Ser Thr Asn Tyr Arg Leu  
 145 150 155  
 ATA AGT TGT AAT ACC TCA GTC ATT ACA CAG GCC TGT CCA 505.  
 Ile Ser Cys Asn Thr Ser Val Ile Thr Gln Ala Cys Pro  
 160 165  
 25 AAG GTA TCC TTT GAG CTA ATT CCC ATA CAT TAT TGT GCC 544  
 Lys Val Ser Phe Glu Leu Ile Pro Ile His Tyr Cys Ala  
 170 175 180  
 CCG GCT GGT TTT GCG ATT CTA AAG TGT AAA GAT AAG AAG 583  
 Pro Ala Gly Ile Leu Lys Cys Lys Asp Lys Lys  
 30 185 190  
 TTC AAT GGA ACA GGA CCA TGT AAA AAT GTC AGC ACA GTA 622  
 Phe Asn Gly Thr Gly Pro Cys Lys Asn Val Ser Thr Val  
 195 200 205  
 CAA TGT ACA CAT GGA ATT AGA CCA CTA CTA TCA ACT CAA 661  
 35 Gln Cys Thr His Gly Ile Arg Pro Val Val Ser Thr Gln  
 210 215 220  
 CTA CTG TTA AAT GGC AGT CTA GCA GAA GAA GAG ATA GTA 700.  
 Leu Leu Leu Asn Gly Ser Leu Ala Glu Glu Ile Val  
 225 230  
 40 ATT AGA TCT GAA AAT ATC ACA GAC AAT GCT AAA ACC ATA 739  
 Ile Arg Ser Glu Asn Ile Thr Asp Asn Ala Lys Thr Ile  
 235 240 245  
 ATA GTG CAG CTA AAT GAA TCT ATA GTG ATT AAT TGT ACA 778  
 Ile Val Gln Leu Asn Glu Ser Ile Val Ile Asn Cys Thr  
 45 250 255  
 AGA CCC AAT AAC AAC ACA AGA AAA ACT ATA AAT ATA GGA 817  
 Arg Pro Asn Asn Asn Thr Arg Lys Ser Ile Asn Ile Gly  
 260 265 270  
 CCA GGG AGA GCA TTC TAT ACA ACA GGA GAC ATA ATA GGA 856  
 50 Pro Gly Arg Ala Phe Tyr Thr Gly Asp Ile Ile Gly  
 275 280 285  
 GAT ATA AGA CAA GCA CAT TGT AAC CTT AGT AAA ACA CAA 895.  
 Asp Ile Arg Gln Ala His Cys Asn Leu Ser Lys Thr Gln  
 290 295  
 55 TGG GAA AAA ACG TTA AGA CAG ATA GCT ATA AAA TTA GAA 934  
 Trp Glu Lys Thr Leu Arg Gln Ile Ala Ile Lys Leu Glu  
 300 305 310  
 GAA AAA TTT AAG AAT AAA ACA ATA CCC TTT AAT AAA TCC 973  
 Glu Lys Phe Lys Asn Lys Thr Ile Ala Phe Asn Lys Ser  
 60 315 320  
 TCA GGA GGG GAC CCA GAA ATT CTA ATC CAC AGT TTT AAT 1012  
 Ser Gly Gly Asp Pro Glu Ile Val Met His Ser Phe Asn  
 325 330 335

TGT GGA GGG GAA TTT TTC TAC TGT AAT ACA ACA AAA CTG 1051  
 Cys Gly Glu Phe Phe Tyr Cys Asn Thr Thr Lys Leu  
 340 345 350  
 TTT AAT ACT ACC TCG AAT TTA ACA CAA CCG TTT AGT AAT 1090  
 5 Phe Asn Ser Thr Trp Asn Leu Thr Gln Pro Phe Ser Asn  
 355 360  
 ACC GGG AAT CGT ACT GAA GAG TTA AAT ATT ACA CTC CCA 1129  
 Thr Gly Asn Arg Thr Glu Glu Leu Asn Ile Thr Leu Pro  
 365 370 375  
 10 TGC AGA ATA AAA CAA ATC ATA AAC TTG TGG CAG GAA GTA 1168  
 Cys Arg Ile Lys Gln Ile Ile Asn Leu Trp Gln Glu Val  
 380 385  
 GGC AAA GCA ATG TAT GCC CCT CCC ATC AGA GGA CAA ATT 1207  
 Gly Lys Ala Met Tyr Ala Pro Pro Ile Arg Gly Gln Ile  
 15 390 395 400  
 AGA TGT TCA TCA AAT ATT ACA GGG CTA CTA TTA ACA AGA 1246  
 Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg  
 405 410 415  
 GAT GGT GGA AGT AAC ACC GGT GAC AAC AGG ACT GAG ACC 1285  
 20 Asp Gly Gly Ser Asn Thr Gly Asp Asn Arg Thr Glu Thr  
 420 425  
 TTT AGA CCT GGA GGA GGA GAT ATG AGG GAC AAT TGG AGA 1324  
 Phe Arg Pro Gly Gly Asp Met Arg Asp Asn Trp Arg  
 430 435 440  
 25 AGT GAA TTA TAT AAA TAT AAA GTA GTA AGA ATT GAA CCA 1363  
 Ser Glu Leu Tyr Lys Tyr Lys Val Val Arg Ile Glu Pro  
 445 450  
 TTA CGA GTA GCA CCC ACC CAG GCA AAG AGA AGA GTG GTG 1402  
 Leu Gly Val Ala Pro Thr Gln Ala Lys Arg Arg Val Val  
 30 455 460 465  
 CAA AGA GAA AAA AGA GCA GTG GGG ATA CGA GCT ATG TTC 1441  
 Gln Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Met Phe  
 470 475 480  
 CTT GGG TTC TTG GGA GAT AA 1461  
 35 Leu Gly Phe Leu Gly Asp  
 485 486

CLONE C8.6

G GTA CCT GTG TGG AAA GAA GCA ACC ACC ACT CTA TTT 37  
 Val Pro Val Trp Lys Glu Ala Thr Thr Thr Leu Phe  
 40 1 5 10  
 TGT GCA TCA GAT GCT AAA CCA TAT GAT ACA GAG GTA CAT 76  
 Cys Ala Ser Asp Ala Lys Tyr Asp Thr Glu Val His  
 15 20 25  
 AAT GTT TGG GCT ACA CAT GCC TGT GTA CCC ACA GAC CCC 115  
 Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro  
 45 30 35  
 AAC CCA CAA GAA GTA CTA TTG GAA AAT GTA ACA GAA AAT 154  
 Asn Pro Gln Glu Val Val Leu Glu Asn Val Thr Glu Asn  
 50 40 45 50  
 TTT AAC ATG TGG AAA AAT AAC ATG GTA GAA CAG ATG CAT 193  
 Phe Asn Met Trp Lys Asn Asn Met Val Glu Gln Met His  
 55 55 60  
 GAG GAT ATA ATC AGT TTA TGG GAT CAA AGT CTA AAG CCA 232  
 Glu Asp Ile Ile Ser Leu Trp Asp Gln Ser Leu Lys Pro  
 55 65 70 75  
 TGT GTA AAA TTA ACC CCA CTC TGT GTT ACT TTA AAT TGC 271  
 Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu Asn Cys  
 80 85 90  
 60 ACT AAT TTG GAG AAT GCT AAT AAT ACC GAG AAT GCT AAT 310  
 Thr Asn Leu Glu Asn Ala Asn Asn Thr Glu Asn Ala Asn  
 95 100  
 AAT ACC AAT AAT TAT ACC TTG GGG ATG GAG AGA GGT GAA 349  
 Asn Thr Asn Asn Tyr Thr Leu Gly Met Glu Arg Gly Glu  
 65 105 110 115

AGA AAA AAC TGC TCT TTC AAT ATC ACC ACA AGC TTA AGA 388  
 Arg Lys Asn Cys Ser Phe Asn Ile Thr Thr Ser Leu Arg  
 120 125  
 GAT AAG GGG AAA AAA GAA TAT GCA TTG TTT TAT AAA CTT 427  
 5 Asp Lys Gly Lys Lys Glu Tyr Ala Leu Phe Tyr Lys Leu  
 130 135 140  
 GAT GTA GTA CAA ATA GAT AAT AGT ACC AAC TAT AGG CTG 466  
 Asp Val Val Gln Ile Asp Asn Ser Thr Asn Tyr Arg Leu  
 145 150 155  
 10 ATA AGT TGT AAT ACC TCA GTC ATT ACA CAG CCC TGT CCA 505  
 Ile Ser Cys Asn Thr Ser Val Ile Thr Gln Ala Cys Pro  
 160 165  
 AAG GTA TCC TTT GAG CCA ATT CCC ATA CAT TAT TGT GCC 544  
 Lys Val Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala  
 170 175 180  
 CCG GCT GGT TTT GCG ATT CTA AAG TGT AAA GAT AAG AAG 583  
 Pro Ala Gly Phe Ala Ile Leu Lys Cys Lys Asp Lys Lys  
 185 190  
 TTC AAT GGA ACA GGA CCA TGT AAA AAT GTC AGG ACA GTA 622  
 Phe Asn Gly Thr Gly Pro Cys Lys Asn Val Arg Thr Val  
 195 200 205  
 CAA TGT ACA CAT GGA ATT AGA CCA GTA GTA TCA ACT CAA 661  
 Gln Cys Thr His Gly Ile Arg Pro Val Val Ser Thr Gln  
 210 215 220  
 25 CTA CTG TTA AAT GGC AGT CTA GCA GAA GAA GAG ATA GTA 700  
 Leu Leu Leu Asn Gly Ser Leu Ala Glu Glu Glu Ile Val  
 225 230  
 ATT AGA TCT GAA AAT ATC ACA GAC AAT GCT AAA ACC ATA 739  
 Ile Arg Ser Glu Asn Ile Thr Asp Asn Ala Lys Thr Ile  
 235 240 245  
 ATA GTG CAG CTA AAT GAA TCT ATA GTG ATT AAT TGT ACA 778  
 Ile Val Gln Leu Asn Glu Ser Ile Val Ile Asn Cys Thr  
 250 255  
 AGA CCC AAT AAC AAC ACA AGA AAA AGT ATA AAT ATA GGA 817  
 35 Arg Pro Asn Asn Asn Thr Arg Lys Ser Ile Asn Ile Gly  
 260 265 270  
 CCA GGG AGA GCA TTC TAT ACA ACA GGA GAC ATA ATA GGA 856  
 Pro Gly Arg Ala Phe Tyr Thr Gly Asp Ile Ile Gly  
 275 280 285  
 40 GAT ATA AGA CAA GCA CAT TGT AAC CTT AGT AAA ACA CAA 895  
 Asp Ile Arg Gln Ala His Cys Asn Leu Ser Lys Thr Gln  
 290 295  
 TGG GAA AAA ACG TTA AGA CAG ATA GCT ATA AAA TTA GAA 934  
 Trp Glu Lys Thr Leu Arg Gln Ile Ala Ile Lys Leu Glu  
 45 300 305 310  
 GAA AAA TTT AAG AAT AAA ACA ATA GCC TTT AAT AAA TCC 973  
 Glu Lys Phe Lys Asn Lys Thr Ile Ala Phe Asn Lys Ser  
 315 320  
 TCA GGA GGG GAC CCA GAA ATT GTA ATG CAC AGT TTT AAT 1012  
 50 Ser Gly Gly Asp Pro Glu Ile Val Met His Ser Phe Asn  
 325 330 335  
 TGT GGA GGG GGA TTT TTC TAC TGT AGT ACG AGA AAA CTG 1051  
 Cys Gly Gly Gly Phe Phe Tyr Cys Ser Thr Arg Lys Leu  
 340 345 350  
 55 TTT AAT AGT ACC TGG AAT TTA ACA CAA CCG TTT AGT AAT 1090  
 Phe Asn Ser Thr Trp Asn Leu Thr Gln Pro Phe Ser Asn  
 355 360  
 ACC GGG GAT CGT ACT GAA GAG TTA AAT ATT ACA CTC CCA 1129  
 Thr Gly Asp Arg Thr Glu Glu Leu Asn Ile Thr Leu Pro  
 60 365 370 375  
 TGC AGA ATA AAA CAA ATC ATA AAC TTG TGG CAG GAA CTA 1168  
 Cys Arg Ile Lys Gln Ile Ile Asn Leu Trp Gln Glu Val  
 380 385

390 GGC AAA GCA ATG TAT GCC CCT CCC ATC AGA GGA CAA ATT 1207  
 Gly Lys Ala Met Tyr Ala Pro Pro Ile Arg Gly Gln Ile  
 395 395 400  
 5 AGA TGT TCA TCA AAT ATT ACA GGG CTA CTA TTA AGG AGA 1246  
 Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Arg Arg  
 405 410 415  
 GAT GGT GGA AGT AAC ACC AGT GAC AAC CAG ACT GAG ACC 1285  
 Asp Gly Gly Ser Asn Thr Ser Asp Asn Gln Thr Glu Thr  
 420 425  
 10 TTT AGA CCT GGG GGA GGA GAT ATG AGG GAC AAG TGG AGA 1324  
 Phe Arg Pro Gly Gly Asp Met Arg Asp Lys Trp Arg  
 430 435 440  
 AGT GAA TTA TAT AAA TAT AAA GTA GTA AGA ATT GAA CCA 1363  
 Ser Glu Leu Tyr Lys Tyr Lys Val Val Arg Ile Glu Pro  
 445 450  
 15 TTA GGA GTA GCA CCC ACC CAG GCA AAG AGA AGA GTG GTG 1402  
 Leu Gly Val Ala Pro Thr Gln Ala Lys Arg Arg Val Val  
 455 460 465  
 20 CAA AGA GAA AAA AGA GCA GTG GGG ATA GGA GCT ATG TTC 1441  
 Gln Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Met Phe  
 470 475 480  
 CTT AGG TTC TTA GGA GAT AAA GCT TCT AGA GTC 1474  
 Leu Arg Phe Leu Gly Asp Lys Ala Ser Arg Val  
 485 490 491  
 25 **CLONE C15.2**  
 1 CTC GAG GTA CCT GTC TCG AAA GAA GCA ACT ACC ACT 36  
 Leu Glu Val Pro Val Trp Lys Glu Ala Thr Thr Thr  
 1 5 10  
 30 CTA TTT TGT GCA TCA GAT GCT AAA GCA TAT AAT ACA GAG 75  
 Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asn Thr Glu  
 15 20 25  
 AAA CAT AAT GTT TGG GCC ACA CAC GCC TGT GTA CCC ACA 114  
 Lys His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr  
 35 30 35  
 GAT CCC AAC CCA CAA GAA GTA GTC TTG GGA AAT GTG ACA 153  
 Asp Pro Asn Pro Gln Glu Val Val Leu Gly Asn Val Thr  
 40 45 50  
 GAA AAT TTT AAC ATG TGG AAA AAT AAC ATG GTA GAA CAA 192  
 Glu Asn Phe Asn Met Trp Lys Asn Asn Met Val Glu Gln  
 55 55 60  
 ATG CAT GAA GAT ATA ATC AGT TTA TGG GAT CAA AGT CTA 231  
 Met His Glu Asp Ile Ile Ser Leu Trp Asp Gln Ser Leu  
 65 70 75  
 45 AAG CCA TGT GTA AAA TTA ACC CCA CTC TGT GTT ACT TTA 270  
 Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu  
 80 85 90  
 AAT TGC ACT GAT GAT TTA GGG AAT GCT ACT AAT ACC AAT 309  
 Asn Cys Thr Asp Asp Leu Gly Asn Ala Thr Asn Thr Asn  
 50 95 100  
 AGT AGT GCC ACT ACC AAT AGT AGT AGT TGG GAA GAA ATG 348  
 Ser Ser Ala Thr Thr Asn Ser Ser Ser Trp Glu Glu Met  
 105 110 115  
 55 AAG GGG GAA ATG AAA AGA TGC TCT TTC AAT ATC ACC ACA 387  
 Lys Gly Glu Met Lys Arg Cys Ser Phe Asn Ile Thr Thr  
 120 125  
 AGC ATA AGA GAT AAG ATT AAG AAA GAA CAT GCA CTT TTC 426  
 Ser Ile Arg Asp Lys Ile Lys Lys Glu His Ala Leu Phe  
 130 135 140  
 60 TAT AGA CTT GAT GTA GTC CCA ATA GAT AAT GAT AAT ACC 465  
 Tyr Arg Leu Asp Val Val Pro Ile Asp Asn Asp Asn Thr  
 145 150 155

ACA TAT AGG TTG ATA AAT TGT AAT ACC TCA GTC ATT ACA 504  
 Thr Tyr Arg Leu Ile Asn Cys Asn Thr Ser Val Ile Thr  
 160 165  
 CAG GCC TCT CCA AAG GTA TCA TTT GAC CCA ATT CCC ATA 543  
 5 Gln Ala Cys Pro Lys Val Ser Phe Glu Pro Ile Pro Ile  
 170 175 180  
 CAT TTT TGT GCC CCG GCT GGT TTT GCG ATT CTA AAG TGT 582  
 His Phe Cys Ala Pro Ala Gly Phe Ala Ile Leu Lys Cys  
 185 190  
 10 AAT AAT AAG ACG TTC GAG GGA AAA GGA CCA TGT AAA AAT 621  
 Asn Asn Lys Thr Phe Glu Gly Lys Gly Pro Cys Lys Asn  
 195 200 205  
 GTC AGT ACA GTA CAA TGC ACA CAT GGA ATT AGG CCA GTA 660  
 Val Ser Thr Val Gln Cys Thr His Gly Ile Arg Pro Val  
 15 210 215 220  
 GTG TCA ACT CAA CTG CTG TTA AAT GGC AGT CTA GCA GAA 699  
 Val Ser Thr Gln Leu Leu Asn Gly Ser Leu Ala Glu  
 225 230  
 GAA GAG GTA ATA ATT AGA TCT GAC AAT ATC ACA GAC AAT 738  
 20 Glu Glu Val Ile Ile Arg Ser Asp Asn Ile Thr Asp Asn  
 235 240 245  
 ACT AAA ACC ATT ATA GTA CAG CTA AAC GAA TCT GTA GTA 777  
 Thr Lys Thr Ile Ile Val Gln Leu Asn Glu Ser Val Val  
 250 255  
 25 ATT AAT TGT ACA AGA CCC AAC AAC AAT ACA AGA AAA AGT 816  
 Ile Asn Cys Thr Arg Pro Asn Asn Asn Thr Arg Lys Ser  
 260 265 270  
 ATA CAT ATA GGA CCA GGG AGT GCA TTT TTT GCA ACA GGA 855  
 Ile His Ile Gly Pro Gly Ser Ala Phe Phe Ala Thr Gly  
 30 275 280 285  
 GAA ATA ATA CGA GAT ATA AGA CAA GCA CAC TGT AAC CTT 894  
 Glu Ile Ile Gly Asp Ile Arg Gln Ala His Cys Asn Leu  
 290 295  
 AGT AGA ACA CAA TGG AAT AAC ACT TTA GGA AAG ATA GTC 933  
 35 Ser Arg Thr Gln Trp Asn Asn Thr Leu Gly Lys Ile Val  
 300 305 310  
 ATA AAA TTA AGA GAA CAA TTT AGA AAA CAA TTT GGA GAA 972  
 Ile Lys Leu Arg Glu Gln Phe Arg Lys Gln Phe Gly Glu  
 315 320  
 40 AAA ACA ATA GTC TTT AAT CGA TCC TCA GGA GGG GAC CCG 1011  
 Lys Thr Ile Val Phe Asn Arg Ser Ser Gly Gly Asp Pro  
 325 330 335  
 GAA ATT GCA ATG CAC AGT TTT AAT TGT GGA GGG GAA TTT 1050  
 Glu Ile Ala Met His Ser Phe Asn Cys Gly Gly Glu Phe  
 45 340 345 350  
 TTC TAC TGT AAC ACA ACA GCA CTG TTT AAT AGT ACC TGG 1089  
 Phe Tyr Cys Asn Thr Thr Ala Leu Phe Asn Ser Thr Trp  
 355 360  
 AAT GTT ACT AAA GGG TTG AAT AAC ACT GAA GGA AAT AGC 1128  
 50 Asn Val Thr Lys Gly Leu Asn Asn Thr Glu Gly Asn Ser  
 365 370 375  
 ACA GGA GAT GAA AAT ATC ATA CTC CCA TGT AGA ATA AAA 1167  
 Thr Gly Asp Glu Asn Ile Ile Leu Pro Cys Arg Ile Lys  
 380 385  
 55 CAA ATT ATA AAC ATG TGG CAG GAA GTA GGA AAA GCA ATG 1206  
 Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys Ala Met  
 390 395 400  
 TAT GCC CCT CCC ATC AGT GGA CAA ATT AGA TGT TCA TCA 1245  
 Tyr Ala Pro Pro Ile Ser Gly Gln Ile Arg Cys Ser Ser  
 60 405 410 415  
 AAC ATT ACA GGG CTG CTA CTA ACA AGA GAT GGT GGT AGT 1284  
 Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly Ser  
 420 425

AAG AAC GAG AGC ATC ACC ACC GAG GTC TTC AGA CCT GGA 1323  
 Lys Asn Glu Ser Ile Thr Thr Glu Val Phe Arg Pro Gly  
 430 435 440  
 CGA CGA GAT ATG AGG GAC AAT TGG AGA AGT GAA TTA TAT 1362  
 5 Gly Gly Asp Met Arg Asp Asn Trp Arg Ser Glu Leu Tyr  
 445 450  
 AAA TAT AAA GTA GTA AAA ATT GAA CCA TTA GGA GTA GCG 1401  
 Lys Tyr Lys Val Val Lys Ile Glu Pro Leu Gly Val Ala  
 455 460 465  
 10 CCC ACC AAG GCA AAG AGA AGA GTG GTG CAG AGA GAA AAA 1440  
 Pro Thr Lys Ala Lys Arg Arg Val Val Gln Arg Glu Lys  
 470 475 480  
 AGA GCA GTG GGA ACA ATA GGA GCT ATG TTC CTT GGG TTC 1479  
 Arg Ala Val Gly Thr Ile Gly Ala Met Phe Leu Gly Phe  
 15 485 490  
 TTC GGA GCA TAA AGC TTC TAG AGT CGA CCT GCA 1512  
 Leu Gly Ala Xaa Ser Phe Xaa Ser Arg Pro Ala  
 495 500 504

CLONE C15.3  
 20 CTC GAG GTA CCT GTG TGG AAA GAA GCA ACT ACC ACT 36  
 Leu Glu Val Pro Val Trp Lys Glu Ala Thr Thr Thr  
 1 5 10  
 CTA TTT TGT GCA TCA GAT GCT AAA GCA TAT AAT ACA GAG 75  
 25 Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asn Thr Glu  
 15 20 25  
 AAA CAT AAT GTT TGG GCC ACA CAC GCC TGT GTC CCC ACA 114  
 Lys His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr  
 30 35  
 GAT CCC AAC CCA CAA GAA GTA GTA TTG GGA AAT GTC ACA 153  
 Asp Pro Asn Pro Gln Glu Val Val Leu Gly Asn Val Thr  
 40 45 50  
 GAA AAT TTT AAC ATG TGG AAA AAT AAC ATG GTA CAA CAA 192  
 Glu Asn Phe Asn Met Trp Lys Asn Asn Met Val Glu Cln  
 35 55 60  
 ATG CAT GAA GAT ATA ATC AGT TTA TGG GAT CAA AGT CTA 231  
 Met His Glu Asp Ile Ile Ser Leu Trp Asp Cln Ser Leu  
 65 70 75  
 AAG CCA TGT GTA AAA TTA ACC CCA CTC TGT GTT ACT TTA 270  
 40 Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu  
 80 85 90  
 AAT TGC ACT GAT GAT TTA GGG AAT GCT ACT AAT ACC AAT 309  
 Asn Cys Thr Asp Asp Leu Gly Asn Ala Thr Asn Thr Asn  
 95 100  
 45 AGC AGT GCC ACT ACC AAT AGT AGT AGT TGG GAA GAA ATG 348  
 Ser Ser Ala Thr Thr Asn Ser Ser Ser Trp Glu Glu Met  
 105 110 115  
 AAG GGG GAA ATG AAA AGG TGC TCT TTC AAT ATC ACC ACA 387  
 Lys Gly Glu Met Lys Arg Cys Ser Phe Asn Ile Thr Thr  
 50 120 125  
 AGC ATA AGA GAT AAG ATT AAG AAA GAA CAT GCA CTT TTC 426  
 Ser Ile Arg Asp Lys Ile Lys Lys Glu His Ala Leu Phe  
 130 135 140  
 TAT AGA CTT GAT GTA GTA CCA ATA GAT AAT GAT AAT ACC 465  
 55 Tyr Arg Leu Asp Val Val Pro Ile Asp Asn Asp Asn Thr  
 145 150 155  
 ACA TAT AGG TTG ATA AAT TGT AAT ACC TCA GTC ATT ACA 504  
 Thr Tyr Arg Leu Ile Asn Cys Asn Thr Ser Val Ile Thr  
 160 165  
 60 CAG GCC TGT CCA AAG GTA TCA TTT GAG CCA ATT CCC ATA 543  
 Gln Ala Cys Pro Lys Val Ser Phe Glu Pro Ile Pro Ile  
 170 175 180  
 CAT TTT TGT GCC CCG GCT GGT TTT GCG ATT CTA AAG TGT 582  
 His Phe Cys Ala Pro Ala Gly Phe Ala Ile Leu Lys Cys  
 65 185 190

	AAT	AAT	AAG	ACG	TTC	GAG	CCA	AAA	CGA	CCA	TGT	AAA	AAT	621
	Asn	Asn	Lys	Thr	Phe	Glu	Gly	Lys	Gly	Pro	Cys	Lys	Asn	
195					200					205				
5	GTC	ACT	ACA	GTA	CAA	TGC	ACA	CAT	GGA	ATT	AGG	CCA	GTA	660
	Val	Ser	Thr	Val	Gln	Cys	Thr	His	Gly	Ile	Arg	Pro	Val	
					210				215		220			
	GTG	TCA	ACT	CAA	CTG	CTG	TTA	AAT	GGC	AGT	CTA	GCA	GAA	699
	Val	Ser	Thr	Gln	Leu	Leu	Asn	Gly	Ser	Leu	Ala	Glu		
					225				230		230			
10	GAA	GAG	GTA	ATA	ATT	AGA	TCT	GCC	AAT	ATC	ACA	GAC	AAT	738
	Glu	Glu	Val	Ile	Ile	Arg	Ser	Gly	Asn	Ile	Thr	Asp	Asn	
					235			240		245				
	ACT	AAA	ACC	ATT	ATA	GTA	CAG	CTA	AAC	GAA	TCT	GTA	GTA	777
	Thr	Lys	Thr	Ile	Ile	Val	Gln	Leu	Asn	Glu	Ser	Val	Val	
15					250			255						
	ATT	AAT	TGT	ACA	AGA	TCC	AAC	AAC	AAT	ACA	AGA	AAA	AGT	816
	Ile	Asn	Cys	Thr	Arg	Ser	Asn	Asn	Asn	Asn	Thr	Arg	Lys	Ser
					260			265		270				
20	ATA	CAT	ATA	GGA	CCA	GGG	AGT	GCA	TTT	TTT	GCA	ACA	GGA	855
	Ile	His	Ile	Gly	Pro	Gly	Ser	Ala	Phe	Phe	Ala	Thr	Gly	
					275			280		285				
	GAA	ATA	ATA	GGA	GAT	ATA	AGA	CAA	GCA	CAC	TGT	AAC	CTT	894
	Glu	Ile	Ile	Gly	Asp	Ile	Arg	Gln	Ala	Ala	His	Cys	Asn	Leu
					290			295						
25	AGT	AGA	ACA	CAA	TGG	AAT	AAC	ACT	TTA	GGA	AAG	ATA	GTC	933
	Ser	Arg	Thr	Gln	Trp	Asn	Asn	Asn	Thr	Leu	Gly	Ile	Val	
					300			305		310				
	ATA	AAA	TTA	AGA	GAA	CAA	TTT	AGA	AAA	CAA	TTT	GGA	GAA	972
	Ile	Lys	Leu	Arg	Glu	Gln	Phe	Arg	Lys	Gln	Phe	Gly	Glu	
30					315			320						
	AAA	ACA	ATA	GTC	TTT	AAT	CGA	TCC	TCA	GGA	GGG	GAC	CCG	1011
	Lys	Thr	Ile	Val	Phe	Asn	Arg	Ser	Ser	Gly	Gly	Asp	Pro	
					325			330		335				
35	GAA	ATT	GCA	ATG	CAC	AGT	TTT	AAT	TGT	GGA	GGG	GAA	TTT	1050
	Glu	Ile	Ala	Met	His	Ser	Phe	Asn	Cys	Gly	Gly	Glu	Phe	
					340			345		350				
	TTC	TAC	TGT	AAC	ACA	ACA	GCA	CTG	TTT	AAT	AGT	ACC	TGG	1089
	Phe	Tyr	Cys	Asn	Thr	Thr	Ala	Leu	Phe	Asn	Ser	Thr	Trp	
40	AAT	GTT	ACT	AAA	GGG	TTG	AAT	AAC	ACT	GAA	GGA	AAT	AGC	1128
	Asn	Val	Thr	Lys	Gly	Leu	Asn	Asn	Asn	Thr	Glu	Gly	Asn	Ser
					365			370		375				
	ACA	GGG	GAT	GAA	AAT	ATC	ATA	CTC	CCA	TGT	AGA	ATA	AAA	1167
	Thr	Gly	Asp	Glu	Asn	Ile	Ile	Leu	Pro	Cys	Arg	Ile	Lys	
45					380			385						
	CAA	ATT	ATA	AAC	ATG	TGG	CAG	GAA	GTA	GGA	AAA	GCA	ATG	1206
	Gln	Ile	Ile	Asn	Met	Trp	Gln	Glu	Val	Gly	Lys	Ala	Met	
					390			395		400				
50	TAT	GCC	CCT	CCC	ATC	AGT	GGA	CAA	ATT	AGA	TGT	TCA	TCA	1245
	Tyr	Ala	Pro	Pro	Ile	Ser	Gly	Gln	Ile	Arg	Cys	Ser	Ser	
					405			410		415				
	AAT	ATT	ACA	GGG	CTG	CTA	CTA	ACA	AGA	GAT	GGT	GCT	AGT	1284
	Asn	Ile	Thr	Gly	Leu	Leu	Leu	Thr	Arg	Asp	Gly	Gly	Ser	
55					420			425						
	AAG	AAC	GAG	AGC	ATC	ACC	ACC	GAG	GTC	TTC	AGA	CCT	GGA	1323
	Lys	Asn	Glu	Ser	Ile	Thr	Thr	Glu	Val	Phe	Arg	Pro	Gly	
					430			435		440				
	GGA	GGA	GAT	ATG	AGG	GAC	AAT	TGG	AGA	AGT	GAA	TTA	TAT	1362
	Gly	Gly	Asp	Met	Arg	Asp	Asn	Trp	Arg	Ser	Glu	Leu	Tyr	
60					445			450						
	AAA	TAT	AAA	GTA	GTA	AAA	ATT	GAA	CCA	TTA	GGA	GTA	GCG	1401
	Lys	Tyr	Lys	Val	Val	Lys	Ile	Glu	Pro	Leu	Gly	Val	Ala	
					455			460		465				

CCC ACC AAG GCA AAG AGA AGA GTG GTG CAG AGA GAA AAA 1440  
 Pro Thr Lys Ala Lys Arg Arg Val Val Gln Arg Glu Lys  
 470 475 480  
 5. AGA GCA GTG GGA ACA ATA GCA GCT ATG TTC CTT GGG TTC 1479  
 Arg Ala Val Gly Thr Ile Gly Ala Met Phe Leu Gly Phe  
 485 490  
 TTA GGA GCA TAA AGC TTC TAG A 1501  
 Leu Gly Ala Xaa Ser Phe Xaa  
 495 500

10

CLONE C7.2

GG GAA TTC GGA TCC GGG GTA CCT GTG TGG AAG GAA GCA 38  
 Glu Phe Gly Ser Gly Val Pro Val Trp Lys Glu Ala  
 1 5 10  
 15 ACC ACC ACT CTA TTC TGT GCA TCA GAT GCT AGA GCA TAT 77  
 Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Arg Ala Tyr  
 15 20 25  
 GAC ACA GAG GTA CAT AAT GTT TGG GCC ACA CAT GCC TGT 116  
 Asp Thr Glu Val His Asn Val Trp Ala Thr His Ala Cys  
 20 30 35  
 GTA CCC ACA GAC CCT ACT CCA CAA GAA GTA GTT TTG GAA 155  
 Val Pro Thr Asp Pro Ser Pro Gln Glu Val Val Leu Glu  
 40 45 50  
 AAT GTG ACA GAA AAT TTT AAC ATG TGG AAA AAT AAC ATG 194  
 Asn Val Thr Glu Asn Phe Asn Met Trp Lys Asn Asn Met  
 55 60  
 500  
 25 GTA GAA CAA ATG CAT GAG GAT ATA ATT AGT TTA TGG GAT 233  
 Val Glu Gln Met His Asp Ile Ile Ser Leu Trp Asp  
 65 70 75  
 30 CAA AGC TTA AAG CCA TGT GTA AAA TTA ACC CCA CTC TGT 272  
 Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys  
 80 85 90  
 GTT ACT TTA AAT TGC AGT GAT TAT AGG AAT GCT ACT GAT 311  
 Val Thr Leu Asn Cys Ser Asp Tyr Arg Asn Ala Thr Asp  
 35 95 100  
 TAT AAG AAT GCT ACT GAT ACC ACT AGT AGT AAC GAG GGA 350  
 Tyr Lys Asn Ala Thr Asp Thr Ser Ser Asn Glu Gly  
 105 110 115  
 40 AAG ATG GAG AGA GGA GAA ATA AAA AAC TGC TCT TTC AAT 389  
 Lys Met Glu Arg Gly Glu Ile Lys Asn Cys Ser Phe Asn  
 120 125  
 ATT ACC ACA AGC ATA AAA AAT AAG ATG CAG AAA GAA TAT 428  
 Ile Thr Thr Ser Ile Lys Asn Lys Met Gln Lys Glu Tyr  
 130 135 140  
 45 GCA CTT TTC TAT AAA CTT GAT ATA GTA CCA ATA GAT AAT 467  
 Ala Leu Phe Tyr Lys Leu Asp Ile Val Pro Ile Asp Asn  
 145 150 155  
 ACA AGC TAT ACA TTG ATA AGT TGT AAC ACC TCA GTC ATT 506  
 Thr Ser Tyr Thr Leu Ile Ser Cys Asn Thr Ser Val Ile  
 50 160 165  
 ACA CAG GCC TGT CCA AAG GTA TCC TTT GAA CCA ACT CCC 545  
 Thr Gln Ala Cys Pro Lys Val Ser Phe Glu Pro Thr Pro  
 170 175 180  
 55 ATA CAT TAT TGT GCT CCG GCT GGT TTT GCG ATT CTA AAG 584  
 Ile His Tyr Cys Ala Pro Ala Gly Phe Ala Ile Leu Lys  
 185 190  
 TGT AAT GAT AAG AAG TTC AGT GGA AAA GGA GAA TGT AAA 623  
 Cys Asn Asp Lys Lys Phe Ser Gly Lys Gly Glu Cys Lys  
 195 200 205  
 60 AAT GTC AGC ACA GTA CAA TGT ACA CAT GGA ATT AGG CCA 662  
 Asn Val Ser Thr Val Gln Cys Thr His Gly Ile Arg Pro  
 210 215 220  
 GTA GTA TCA ACT CAA CTG CTG TTA AAT GGC AGT CTA GCA 701  
 Val Val Ser Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala  
 225 230

GAA GAA GAG GTG GTA ATT AGA TCT GAC AAT TTC ATA GAC 740  
 Glu Glu Val Val Ile Arg Ser Asp Asn Phe Ile Asp  
 235 240 245  
 5 AAT ACT AAA ACC ATA ATA CTA CAG CTG AAA GAA TCT GTA 779  
 Asn Thr Lys Thr Ile Ile Val Gln Leu Lys Glu Ser Val  
 250 255  
 GAA ATT AAT TGT ATA AGA CCC AAC AAT AAT ACA AGA AAA 818  
 Glu Ile Asn Cys Ile Arg Pro Asn Asn Thr Arg Lys  
 260 265 270  
 10 GGT ATA CAT ATA GGA CCA GGG AGA GCA TGG TAT GCA ACA 857  
 Gly Ile His Ile Gly Pro Gly Arg Ala Trp Tyr Ala Thr  
 275 280 285  
 GGA GAA ATA GTG GGA GAT ATA AGA AAG GCA TAT TGT AAC 896  
 Gly Glu Ile Val Gly Asp Ile Arg Lys Ala Tyr Cys Asn  
 15 290 295  
 ATT AGT AGA ACA AAA TGG AAT AAC ACT TTA ATA CAG ATA 935  
 Ile Ser Arg Thr Lys Trp Asn Asn Thr Leu Ile Gln Ile  
 300 305 310  
 20 GCT AAC AAA TTA AAA GAA AAA TAT AAT ACA ACA ATA ACC 974  
 Ala Asn Lys Leu Lys Glu Lys Tyr Asn Thr Thr Ile Ser  
 315 320  
 TTT AAT CGA TCC TCA GGA GGG GAC CCA GAA ATT GCA ACG 1013  
 Phe Asn Arg Ser Ser Gly Gly Asp Pro Glu Ile Val Thr  
 325 330 335  
 25 CAT AGT TTT AAT TGT GGA GGG GAG TTT TTC TAC TGT GAT 1052  
 His Ser Phe Asn Cys Gly Gly Glu Phe Phe Tyr Cys Asp  
 340 345 350  
 TCA ACA CAA CTG TTT AAT AGT ACT TGG AAT TTA AAT CGT 1091  
 Ser Thr Gln Leu Phe Asn Ser Thr Trp Asn Leu Asn Gly  
 355 360  
 30 ACT TGG AAT TTT ACT GCA GGG TCA AAT GAA ACT GAA GGC 1130  
 Thr Trp Asn Phe Thr Ala Gly Ser Asn Glu Thr Glu Gly  
 365 370 375  
 35 AAT ATC ACA CTC CCA TGC AGA ATA AAA CAA ATT ATA AAC 1169  
 Asn Ile Thr Leu Pro Cys Arg Ile Lys Gln Ile Ile Asn  
 380 385  
 AGG TGG CAG GAA GTA GGG AAA GCA ATG TAT GCC CCT CCC 1208  
 Arg Trp Gln Glu Val Gly Lys Ala Met Tyr Ala Pro Pro  
 390 395 400  
 40 ATC AGT GGA CAA ATA AAA TGC TCA TCA AAC ATT ACA GGG 1247  
 Ile Ser Gly Gln Ile Lys Cys Ser Ser Asn Ile Thr Gly  
 405 410 415  
 ATG ATA TTA ACA AGG GAT GGT GGT AAC GAG AAC AAT AAT 1286  
 Met Ile Leu Thr Arg Asp Gly Gly Asn Glu Asn Asn Asn  
 45 420 425  
 GAG AGC AGT ACT ACT GAG ACC TTC AGA CCG GGA GGA GGA 1325  
 Glu Ser Ser Thr Thr Glu Thr Phe Arg Pro Gly Gly Gly  
 430 435 440  
 50 GAT ATG AGG AAC AAT TGG AGA AGT GAA TTA TAT AAA TAT 1364  
 Asp Met Arg Asn Asn Trp Arg Ser Glu Leu Tyr Lys Tyr  
 445 450  
 AAA GCA GCA AAA ATT GAA CCA TTA GGA GTA GCA CCC ACC 1403  
 Lys Val Val Lys Ile Glu Pro Leu Gly Val Ala Pro Thr  
 455 460 465  
 55 AAG GCA AAG AGA AGA GTG GTG CAG AGA GAA AAA AGA GCA 1442  
 Lys Ala Lys Arg Arg Val Val Gln Arg Glu Lys Arg Ala  
 470 475 480  
 GTG GGA GCG CTA GGA GCT ATG TTC CTT GGG TTC TTA GGA 1481  
 Val Gly Ala Leu Gly Ala Met Phe Leu Gly Phe Leu Gly  
 60 485 490  
 GCA TAA AGC TTC TAG ACC GAC TCT AGA GGA TCC 1514  
 Ala Xaa Ser Phe Xaa Thr Asp Ser Arg Gly Ser  
 495 500 504

CLONE C7.10

G CTA CCT GTG TGG AAG GAA GCA ACC ACC ACT CTA TTC 37  
 Val Pro Val Trp Lys Glu Ala Thr Thr Thr Leu Phe  
 1 5 10  
 TGT GCA TCA GAT GCT AGA GCA TAT GAC ACA GAG GTA CAT 76  
 Cys Ala Ser Asp Ala Arg Ala Tyr Asp Thr Glu Val His  
 15 20 25  
 AAT GTT TGG GCC ACA CAT GCC TGT GTA CCC ACA GAC CCT 115  
 Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro  
 10 15 20 25 30 35  
 AGT CCA CAA GAA GTA TTT TTG GGA AAT GTG ACA GAA AAT 154  
 Ser Pro Gin Glu Val Phe Leu Gly Asn Val Thr Glu Asn  
 40 45 50  
 TTT AAT ATG TGG AAA AAT AAC ATG GTA GAA CAA ATG TAT 193  
 Phe Asn Met Trp Lys Asn Asn Met Val Glu Gin Met Tyr  
 15 20 25 30 35 40 45 50 55 60  
 GAG GAT ATA ATT AGT TTA TGG GAT CAA AGC TTA AAG CCA 232  
 Glu Asp Ile Ile Ser Leu Trp Asp Gln Ser Leu Lys Pro  
 65 70 75  
 TGT GTA AAA TTA ACC CCA CTC TGT GTT ACT TTA AAT TGC 271  
 Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu Asn Cys  
 80 85 90  
 AGT GAT TAT AGG AAT GCT ACT GAT TAT AAG AAT GCT ACT 310  
 Ser Asp Tyr Arg Asn Ala Thr Asp Tyr Lys Asn Ala Thr  
 25 30 35 40 45 50 55 60 65 70 75 80 85 90 95 100  
 GAT ACC ACT AGT AGT AAC GAG GGA AAG ATG GAG AGA GGA 349  
 Asp Thr Thr Ser Ser Asn Glu Gly Lys Met Glu Arg Gly  
 105 110 115  
 GAA ATA AAA AAC TGC TCT TTC AAT ATC ACC ACA AGC ATA 388  
 Glu Ile Lys Asn Cys Ser Phe Asn Ile Thr Thr Ser Ile  
 120 125  
 AAA AAT AAG ATG CAG AAA GAA TAT GCA CTT TTC TAT AAA 427  
 Lys Asn Lys Met Gln Lys Glu Tyr Ala Leu Phe Tyr Lys  
 130 135 140  
 CTT AAT ATA GTA CCA ATA GAT AAT ACA AGC TAT ACA TTG 466  
 Leu Asn Ile Val Pro Ile Asp Asn Thr Ser Tyr Thr Leu  
 145 150 155  
 ATA AGT TGT AAC ACC TCA GTC ATT ACA CAG CCC TGT CCA 505  
 Ile Ser Cys Asn Thr Ser Val Ile Thr Gln Ala Cys Pro  
 40 45 50 55 60 65 70 75 80 85 90 95 100 105 110 115 120 125 130 135 140 145 150 155 160 165 170 175 180 185 190 195 200 205 210 215 220 225 230 235 240 245 250 255 260 265 270  
 AAG GTA TCC TTT GAA CCA ATT CCC ATA CAT TAT TGT GCT 544  
 Lys Val Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala  
 170 175 180  
 CCG GCT GGT TTT GCG ATT CTA AAG TGT AAT GAT AAG AAG 583  
 Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asp Lys Lys  
 185 190  
 TTC AGT GGA AAA GGA GAA TGT AAA AAT GTC ACC ACA GTA 622  
 Phe Ser Gly Lys Gly Glu Cys Lys Asn Val Ser Thr Val  
 195 200 205  
 CAA TGT ACA CAT GGA ATT AGG CCA GTA GTA TCA ACT CAA 661  
 Gln Cys Thr His Gly Ile Arg Pro Val Val Ser Thr Gln  
 210 215 220  
 CTG CTG TTA AAT GGC AGT CTA GCA GAA GAA GAG GTG GTA 700  
 Leu Leu Leu Asn Gly Ser Leu Ala Glu Glu Val Val  
 225 230  
 ATT AGA TCT GAC AAT TTC ACA GAC AAT ACT AAA ACC ATA 739  
 Ile Arg Ser Asp Asn Phe Thr Asp Asn Thr Lys Thr Ile  
 235 240 245  
 ATA GTA CAG CTG AAA GAA TCT GTA GAA ATT AAT TGT ATA 778  
 60 Ile Val Gln Leu Lys Glu Ser Val Glu Ile Asn Cys Ile  
 250 255  
 AGA CCC AAC AAT AAT ACA AGA AAA GGT ATA CAT ATA GGA 817  
 Arg Pro Asn Asn Asn Thr Arg Lys Gly Ile His Ile Gly  
 260 265 270

CCA GGG AGA GCA TGG TAT GCA ACA GGA GAA ATA GTA GGA 856  
 Pro Gly Arg Ala Trp Tyr Ala Thr Gly Glu Ile Val Gly  
 275 280 285  
 GAT ATA AGA CAG GCA TAT TGT AAC ATT ACT ACA ACA AAA 895  
 5 Asp Ile Arg Gln Ala Tyr Cys Asn Ile Ser Arg Thr Lys  
 290 295  
 TGG AAT AAC ACT TTA ATA CAG ATA GCT AAC AAA TTA AAA 934  
 Trp Asn Asn Thr Leu Ile Gln Ile Ala Asn Lys Leu Lys  
 300 305 310  
 10 GAA AAA TAT AAT ACA ACA ATA AGC TTT AAT CGA CAA TCC TCA 973  
 Glu Lys Tyr Asn Thr Thr Ile Ser Phe Asn Arg Ser Ser  
 315 320  
 GGA GGG GAC CCA GAA ATT GTA ACC CAT AGT TTT AAT TGT 1012  
 Gly Gly Asp Pro Glu Ile Val Thr His Ser Phe Asn Cys  
 15 325 330 335  
 GGA GGG GAA TTT TTC TAC TGT AAT TCA ACA CAA CTG TTT 1051  
 Gly Gly Glu Phe Phe Tyr Cys Asn Ser Thr Gln Leu Phe  
 340 345 350  
 AAT AGT ACT TGG AAT TTA AAT GGT ACT TGG AAT TTT ACT 1090  
 20 Asn Ser Thr Trp Asn Leu Asn Gly Thr Trp Asn Phe Thr  
 355 360  
 GCA GGG TCA AAT GAA ACT GAA GGC AAT ATC ACA CTC CCA 1129  
 Ala Gly Ser Asn Glu Thr Glu Gly Asn Ile Thr Leu Pro  
 365 370 375  
 25 TGC AGA ATA AAA CAA ATT ATA AAC AGG TGG CAG GAA GTC 1158  
 Cys Arg Ile Lys Gln Ile Ile Asn Arg Trp Gln Glu Val  
 380 385  
 GGA AAA GCA ATG TAT GCC CCT CCC ATC AGT GGA CAA ATA 1207  
 Gly Lys Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile  
 390 395 400  
 30 AGA TGC TCA TCA AAC ATT ACA GGG ATG ATA TTA ACA AGG 1246  
 Arg Cys Ser Ser Asn Ile Thr Gly Met Ile Leu Thr Arg  
 405 410 415  
 GAT GGT GGT AAC GAG AAC AAT AAT GAG AGC AGT ACT ACT 1285  
 35 Asp Gly Gly Asn Glu Asn Asn Asn Glu Ser Ser Thr Thr  
 420 425  
 GAG ACC TTC AGA CCG GGA GGA GAT ATG AGG AAC AAT 1324  
 Glu Thr Phe Arg Pro Gly Gly Asp Met Arg Asn Asn  
 430 435 440  
 40 TGG AGA AGT GAA TTA TAT AAA TAT AAA GTA GTA AAA ATT 1363  
 Trp Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile  
 445 450  
 GAG CCA TTA GGA GTA GCA CCC ACC GAC TCT AGA GGA TCC 1402  
 Glu Pro Leu Gly Val Ala Pro Thr Asp Ser Arg Gly Ser  
 45 455 460 465  
 TCT AGA 1408  
 Ser Arg  
 469

50 CLONE C11.5  
 GAG GTA CCT GTG TGG AAA GAA GCA ACC ACT ACT CTA 36.  
 Glu Val Pro Val Trp Lys Glu Ala Thr Thr Leu  
 1 5 10  
 TTT TGT GCA TCA GAT GCT AAA GCA TAT GAC ACA GGG GTG 75.  
 55 Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Gly Val  
 15 20 25  
 CAT AAT GTT TGG GCC ACA CAT GCC TGT GTA CCC ACA GAC 114.  
 His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp  
 30 35  
 60 CCC AAC CCA CAA GAA ATA GAA TTG GTA AAT GTG ACA GAA 153.  
 Pro Asn Pro Gln Glu Ile Glu Leu Val Asn Val Thr Glu  
 40 45 50  
 GAT TTT AAC ATG TGG AAA AAT AAA ATG GTA GAC CAG ATG 192.  
 Asp Phe Asn Met Trp Lys Asn Lys Met Val Asp Gln Met  
 65 55 60

	CAT	GAG	GAT	ATA	ATC	AGT	TTA	TGG	GAT	GAA	AGC	CTA	AAG	231
	His	Glu	Asp	Ile	Ile	Ser	Leu	Trp	Asp	Glu	Ser	Leu	Lys	
	65						70					75		
5	CCA	TGT	GTA	AAG	TTA	ACC	CCA	CTT	TGT	ACT	CTA	AAC	270	
	Pro	Cys	Val	Val	Lys	Leu	Thr	Pro	Leu	Cys	Val	Thr	Leu	Asn
	80						85					90		
	TGC	AGT	GAT	GTG	AAC	AAT	TCC	ACA	AAT	CCT	AAT	GAT	ACT	309
	Cys	Ser	Asp	Val	Asn	Asn	Ser	Thr	Asn	Pro	Asn	Asp	Thr	
	95						100							
10	AAT	ACT	AAT	TCC	ACT	AAT	ACT	ACT	TCC	TCT	ACT	CCT	ACG	348
	Asn	Thr	Asn	Ser	Thr	Asn	Thr	Thr	Ser	Ser	Thr	Pro	Thr	
	105						110					115		
	GCC	ACT	ACT	AGT	AGC	GAG	GAA	AAG	ATG	GAG	AAG	GGA	GAA	387
	Ala	Thr	Thr	Ser	Ser	Glu	Glu	Met	Glu	Lys	Gly	Glu		
15	120						125							
	ATA	AAA	AAC	TGC	TCT	TTC	AAT	ATC	ACC	ACA	CAC	ATG	AAA	426
	Ile	Lys	Asn	Cys	Ser	Phe	Asn	Ile	Thr	Thr	His	Met	Lys	
	130						135					140		
	GAT	AAG	GCA	CAG	AAA	GAA	TAT	GCA	CTT	TTT	TAT	AAA	CTT	465
20	Asp	Lys	Ala	Gln	Lys	Glu	Tyr	Ala	Leu	Phe	Tyr	Lys	Leu	
	145						150					155		
	GAT	ATA	GTA	CCA	ATA	GAT	GAT	AAT	GCC	AGC	TAT	AGG	504	
	Asp	Ile	Val	Pro	Ile	Asp	Asp	Asn	Asn	Ala	Ser	Tyr	Arg	
	160						165							
25	TTG	ATA	AGT	TGT	AAT	ACC	TCA	GAC	ATT	ACA	CAG	GCC	TGT	543
	Leu	Ile	Ser	Cys	Asn	Thr	Ser	Asp	Ile	Thr	Gln	Ala	Cys	
	170						175					180		
	CCA	AAG	GTC	ACC	TTT	GAG	CCA	ATT	CCC	ATA	CAT	TAT	TGT	582
	Pro	Lys	Val	Thr	Phe	Glu	Pro	Ile	Pro	Ile	His	Tyr	Cys	
30	185						190							
	GCC	CCG	GCT	GGT	TTT	GCG	ATT	CTA	AAG	TGT	AAA	GAT	AAG	621
	Ala	Pro	Ala	Gly	Phe	Ala	Ile	Leu	Lys	Cys	Lys	Asp	Lys	
	195						200					205		
	AAG	TTC	AAT	GGA	ACA	GGA	CCA	TGT	TCA	AAG	GTC	AGC	ACA	660
35	Lys	Phe	Asn	Gly	Thr	Gly	Pro	Cys	Ser	Lys	Val	Ser	Thr	
	210						215					220		
	GTA	CAA	TGT	ACA	CAT	GGA	ATT	AGG	CCA	GTA	GTA	TCA	ACT	699
	Val	Gln	Cys	Thr	His	Gly	Ile	Arg	Pro	Val	Val	Ser	Thr	
	225						230							
40	CAA	CTG	TTG	TTA	AAT	GGC	AGT	CTT	GCA	GAA	GAA	GTA	738	
	Gln	Leu	Leu	Leu	Asn	Gly	Ser	Leu	Ala	Glu	Glu	Glu	Val	
	235						240					245		
	GTA	ATT	AGA	TCT	GTC	AAT	TTC	ACA	GAC	AAT	GCT	AAA	ATC	777
	Val	Ile	Arg	Ser	Val	Asn	Phe	Thr	Asp	Asn	Ala	Lys	Ile	
45	250						255							
	ATA	ATA	GTA	CAG	CTG	AAA	GAA	CCT	GTA	GCA	ATT	AAT	TGT	816
	Ile	Ile	Val	Gln	Leu	Lys	Glu	Pro	Val	Ala	Ile	Asn	Cys	
	260						265					270		
	ACA	AGA	CCC	AAC	AAC	AAT	ACA	AGA	AAA	GGT	ATA	CAT	CTA	855
50	Thr	Arg	Pro	Asn	Asn	Asn	Thr	Arg	Lys	Gly	Ile	His	Leu	
	275						280					285		
	GGA	CCA	GGG	AGC	ACA	TTT	TAT	ACA	ACA	GGA	GAA	ATA	ATA	894
	Gly	Pro	Gly	Ser	Thr	Phe	Tyr	Thr	Thr	Gly	Glu	Ile	Ile	
	290						295							
55	GGA	GAC	ATA	AGA	AAA	GCA	TAT	TGC	AAG	ATT	AGT	AAA	GAA	933
	Gly	Asp	Ile	Arg	Lys	Ala	Tyr	Cys	Lys	Ile	Ser	Lys	Glu	
	300						305					310		
	AAA	TGG	AAT	AAC	ACT	TTA	AGA	CAG	GTA	GTT	AAA	AAA	TTA	972
60	Lys	Trp	Asn	Asn	Thr	Leu	Arg	Gln	Val	Val	Lys	Lys	Leu	
	315						320							
	AGA	GAA	CAA	TTT	GGG	AAT	AAA	ACA	ATA	ATT	TTT	AAT	CGA	1011
	Arg	Glu	Gln	Phe	Gly	Asn	Lys	Thr	Ile	Ile	Phe	Asn	Arg	
	325						330					335		

TCC TCA GGA GGG GAC CCA GAA ATT GTA ATG CAC AGT TTT 1050  
 Ser Ser Gly Gly Asp Pro Glu Ile Val Met His S r Phe  
       340                 345                 350  
 AAC TGT CGA GGG GAG TTT TTC TAC TGT AAT ACA ACA CAA 1089  
 Asn Cys Gly Gly Glu Phe Phe Tyr Cys Asn Thr Thr Glu  
       355                 360                 365  
 CTG TTT AAT AGT ACT TGG AAT AAT ACT GAA GGG ACA AAT 1128  
 Leu Phe Asn Ser Thr Trp Asn Asn Thr Glu Gly Thr Asn  
       365                 370                 375  
 10 AGC ACT GAA CGA AAT AGC ACA ATC ACA CTC CCA TGC AGA 1167  
 Ser Thr Glu Gly Asn Ser Thr Ile Thr Leu Pro Cys Arg  
       380                 385                 390  
 ATA AAA CAA ATT ATA AAT ATG TGG CAG GAA GTA GGA AAA 1206  
 Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys  
       390                 395                 400  
 GCA ACG TAT GCC CCT CCC ATC AGA GGA CGA ATT AGA TGC 1245  
 Ala Thr Tyr Ala Pro Pro Ile Arg Gly Arg Ile Arg Cys  
       405                 410                 415  
 ATA TCA AAT ATT ACA GGA CTG CTA TTA ACA AGA GAT GGT 1284  
 Ile Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly  
       420                 425                 430  
 GGT AGG AAT GTC ACA AAC AAT ACC GAA ACC TTC AGA CCT 1323  
 Gly Arg Asn Val Thr Asn Asn Thr Glu Thr Phe Arg Pro  
       430                 435                 440  
 25 GGA GGA GGA GAC ATG AGG GAC AAT TGG AGA AGT GAA TTA 1362  
 Gly Gly Gly Asp Met Arg Asp Asn Trp Arg Ser Glu Leu  
       445                 450                 455  
 TAT AAA TAT AAA GTA GTA AAA GTT GAA CCA TTA GGA ATA 1401  
 Tyr Lys Tyr Lys Val Val Lys Val Glu Pro Leu Gly Ile  
       455                 460                 465  
 GCA CCC ACC AAG GCA AAG AGA AGA GTG GTG CAC AGA GAC 1440  
 Ala Pro Thr Lys Ala Lys Arg Arg Val Val His Arg Asp  
       470                 475                 480  
 AAA AGA GCA GCA CTA GGA GCC TTG TTC CTT GGG TTC TTA 1479  
 Lys Arg Ala Ala Leu Gly Ala Leu Phe Leu Gly Phe Leu  
       485                 490                 495  
 GGA GCA TAA AAG CTT CTA GA 1499  
 Gly Ala Xaa Lys Leu Leu  
       495                 499

CLONE C11.7

GAG GTA CCT GTA TGG AAA GAA GCA ACC ACT ACT CTA 36  
 Glu Val Pro Val Trp Lys Glu Ala Thr Thr Leu  
       1                 5                 10  
 45 TTT TGT GCA TCA GAT GCT AAA GCA TAT GAC ACA GAG GTG 75  
 Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu Val  
       15                 20                 25  
 CAT AAT GTT TGG GCC ACA CAT GCC TGT GTA CCC ACA GAC 114  
 His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp  
       30                 35                 40  
 50 CCC AAC CCA CAA GAA ATA GAA TTG GTA AAT GTG ACA GAA 153  
 Pro Asn Pro Gln Glu Ile Glu Leu Val Asn Val Thr Glu  
       40                 45                 50  
 GAT TTT AAC ATG TGG AAA AAT AAA ATG GTA GAC CAG ATG 192  
 Asp Phe Asn Met Trp Lys Asn Lys Met Val Asp Gln Met  
       55                 60                 65  
 CAT GAG GAT ATA ATC AGT TTA TGG GAT GAA AGC CTA AAG 231  
 His Glu Asp Ile Ile Ser Leu Trp Asp Glu Ser Leu Lys  
       65                 70                 75  
 60 CCA TGT GTA AAG TTA ACC CCA CTT TGT GTT ACT CTA AAC 270  
 Pro Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu Asn  
       80                 85                 90  
 TGC AGT GAT GTG AAC AAT TCC ACA AAT CCT AAT GAT ACT 309  
 Cys Ser Asp Val Asn Asn Ser Thr Asn Pro Asn Asp Thr  
       95                 100

AAT ACT AAT TCC ACT AAT ACT TCC TCT ACT CCT ACG 348  
 Asn Thr Asn Ser Thr Asn Thr Thr Ser Ser Thr Pro Thr  
 105 110 115  
 CCC ACT ACT AGT AGC GAG GAA AAG ATG GAG AAG GGA GAA 387  
 5 Ala Thr Thr Ser Ser Glu Glu Lys Met Glu Lys Gly Glu  
 120 125  
 ATA AAA AAC TGC TCT TTC AAT ATC ACC ACA CAC ATG AAA 426  
 Ile Lys Asn Cys Ser Phe Asn Ile Thr Thr His Met Lys  
 130 135 140  
 10 GAT AAG GTA CAG AAA GAA TAT GCA CTT TTT TAT AAA CTT 465  
 Asp Lys Val Gln Lys Glu Tyr Ala Leu Phe Tyr Lys Leu  
 145 150 155  
 GAT ATA GTA CCA ATA GAT GAT AAT AAT ACC ACC TAT AGG 504  
 Asp Ile Val Pro Ile Asp Asp Asn Asn Thr Ser Tyr Arg  
 15 160 165  
 TTG ATA AGT TGT AAT ACC TCA GTC ATT ACA CAG GCC TGT 543  
 Leu Ile Ser Cys Asn Thr Ser Val Ile Thr Gln Ala Cys  
 170 175 180  
 CCA ATG GTG ACC TTT GAG CCA ATT CCC ATA CAT TAT TGT 582  
 20 Pro Met Val Thr Phe Glu Pro Ile Pro Ile His Tyr Cys  
 185 190  
 GCC CCG CCT GGT TTT GCG ATT CTA AAG TGT AAA GAT AAG 621  
 Ala Pro Ala Gly Phe Ala Ile Leu Lys Cys Lys Asp Lys  
 195 200 205  
 25 AAG TTC AAT GGA ACA GGA CCA TGT TCA AAG GTC AGC ACA 660  
 Lys Phe Asn Gly Thr Gly Pro Cys Ser Lys Val Ser Thr  
 210 215 220  
 GTA CAA TGT ACA CAT GGA ATT AGG CCA GTA GTA TCA ACT 699  
 Val Gln Cys Thr His Gly Ile Arg Pro Val Val Ser Thr  
 30 225 230  
 CAA CTG TTG TTA AAT GGC AGT CTT GCA GAA GAA GAA GTA 738  
 Gln Leu Leu Asn Gly Ser Leu Ala Glu Glu Val  
 235 240 245  
 35 GTA ATT AGA TCT GTC AAT TTC ACA GAC AAT GCT AAA ATC 777  
 Val Ile Arg Ser Val Asn Phe Thr Asp Asn Ala Lys Ile  
 250 255  
 40 ATA ATA GCA CAG CTG AAA GAA CCT GCA GCA ATT AAT TCT 816  
 Ile Ile Val Gln Leu Lys Glu Pro Val Ala Ile Asn Cys  
 260 265 270  
 ACA AGA CCC AAC AAC AAT ACA AGA AAA GGT ATA CAT CTA 855  
 Thr Arg Pro Asn Asn Asn Thr Arg Lys Gly Ile His Leu  
 275 280 285  
 GGA CCA CGG AGC ACA TTT TAT ACA ACA GGA GAA ATA ATA 894  
 Gly Pro Gly Ser Thr Phe Tyr Thr Thr Gly Glu Ile Ile  
 45 290 295  
 GGA GAC ATA AGA AAA GCA TAT TGC AAG ATT AGT AAA GAA 933  
 Gly Asp Ile Arg Lys Ala Tyr Cys Lys Ile Ser Lys Glu  
 300 305 310  
 50 AAA TGG AAT AAC ACT TTA AGA CAG GCA CTT AAA AAA TTA 972  
 Lys Trp Asn Asn Thr Leu Arg Gln Val Val Lys Lys Leu  
 315 320  
 AGA GAA CAA TTT GGG AAT AAA ACA ATA ATT TTT AAT CGA 1011  
 Arg Glu Gln Phe Gly Asn Lys Thr Ile Ile Phe Asn Arg  
 325 330 335  
 55 TCC TCA CGG GGG GAC CCA GAA ATT GCA ATG CAC AGT TTT 1050  
 Ser Ser Gly Gly Asp Pro Glu Ile Val Met His Ser Phe  
 340 345 350  
 AAC TGT CGG GGG GAG TTT TTC TAC TGT AAT ACA ACA CAA 1089  
 Asn Cys Gly Gly Glu Phe Phe Tyr Cys Asn Thr Thr Gln  
 60 355 360  
 CTG TTT AAT AGT ACT TGG AAT AAT ACT GAA GGG ACA AAT 1128  
 Leu Phe Asn Ser Thr Trp Asn Asn Thr Glu Gly Thr Asn  
 365 370 375

AGC ACT GAA GGA AAT AGC ACA ATC ACA CTC CCA TGC AGA 1167  
 Ser Thr Glu Gly Asn Ser Thr Ile Thr Leu Pro Cys Arg  
 380 385  
 ATA AAA CAA ATT ATA AAT ATG TGG CAG GAA GTA GGA AAA 1206  
 5 Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys  
 390 395 400  
 GCA ACG TAT GCC CCT CCC ATC AGA GGA CGA ATT AGA TGC 1245  
 Ala Thr Tyr Ala Pro Pro Ile Arg Gly Arg Ile Arg Cys  
 405 410 415  
 10 ATA TCA AAT ATT ACA GGA CTG CTA TTA ACA AGA GAT GGT 1284  
 Ile Ser Asn Ile Thr Gly Leu Leu Thr Arg Asp Gly  
 420 425  
 GGT AGG AAT GTC ACA AAC AAT ACC GAN NCC TTC AGA CCT 1323  
 Gly Arg Asn Val Thr Asn Asn Thr Xaa Xaa Phe Arg Pro  
 15 430 435 440  
 GGA GGA GGA GAC ATG AGG GAC AAT TGG AGA AGT GAA TTA 1362  
 Gly Gly Gly Asp Met Arg Asp Asn Trp Arg Ser Glu Leu  
 445 450  
 TAT AAA TAT AAA GTA GTA AAA GTT GAA CCA TTA GGA ATA 1401  
 20 Tyr Lys Tyr Lys Val Val Lys Val Glu Pro Leu Gly Ile  
 455 460 465  
 GCA CCC ACC AAG GCA AAG AGA AGA GTG GTG CAC AGA GAC 1440  
 Ala Pro Thr Lys Ala Lys Arg Arg Val Val His Arg Asp  
 470 475 480  
 25 AAA AGA GCA GCA CTA GGA GCT TTG TTC CTT GGG TTC TTA 1479  
 Lys Arg Ala Ala Leu Gly Ala Leu Phe Leu Gly Phe Leu  
 485 490  
 GGA GCA TAA AAG CTT CTA GA 1499  
 Gly Ala Xaa Lys Leu Leu  
 30 495 499

CLONE C10.5

G GTA CCT GTG TGG AAA GAA GCA AAC ACA ACT CTA TTT 37  
 Val Pro Val Trp Lys Glu Ala Asn Thr Thr Leu Phe  
 35 1 5 10  
 TGT GCA TCA GAT GCT AAA GCA TAT GAT AGA GAA GTA CAT 76  
 Cys Ala Ser Asp Ala Lys Ala Tyr Asp Arg Glu Val His  
 15 20 25  
 AAT GTT TGG GCA ACA CAT GCC TGT GTA CCC ACA GAC CCC 115  
 40 Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro  
 30 35  
 AAC CCA CAA GAA ATA GTA TTG GGA AAT GTG ACA GAA AAT 154  
 Asn Pro Gln Glu Ile Val Leu Gly Asn Val Thr Glu Asn  
 40 45 50  
 45 TTT AAC ATG TGG AAA AAT AAC ATG GTA GAA CAA ATG CAT 193  
 Phe Asn Met Trp Lys Asn Asn Met Val Glu Gln Met His  
 55 60  
 GAG GAT ATA ATC AAT TTA TGG GAT CAA AGC TTA AAG CCA 232  
 Glu Asp Ile Ile Asn Leu Trp Asp Gln Ser Leu Lys Pro  
 50 65 70 75  
 TGT GTA AAG TTA ACT CCA CTC TGT GTT ACT TTA AAG TGC 271  
 Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu Lys Cys  
 80 85 90  
 55 AAG GAT CTG GAG AGG AAT ACT ACC TAT AAT AGC ACT ATT 310  
 Lys Asp Leu Glu Arg Asn Thr Thr Tyr Asn Ser Thr Ile  
 95 100  
 ACC AAT AAT AGT AGT TTG GAG GGA CTA AGA GAA CAA ATG 349  
 Thr Asn Asn Ser Ser Leu Glu Gly Leu Arg Glu Gln Met  
 105 110 115  
 60 ACA AAC TGC TCT TTC AAC ATC ACC ACA AGT ATA AGA GAT 388  
 Thr Asn Cys Ser Phe Asn Ile Thr Thr Ser Ile Arg Asp  
 120 125  
 AAG GTG CAG AAA GAA TAT GCA CTT TTG TAT AAA CTT GAT 427  
 Lys Val Gln Lys Glu Tyr Ala Leu Leu Tyr Lys Leu Asp  
 65 130 135 140

GTA GTA CCA ATA GAA GAA GAT GAC AAT ACT AGC TAT AGA 466  
 Val Val Pro Ile Glu Glu Asp Asp Asn Thr Ser Tyr Arg  
 145 150 155  
 TTG ATA AGT TGT AAC ACC TCA GTC ATT ACA CAG GCT TGT 505  
 5 Leu Ile Ser Cys Asn Thr Ser Val Ile Thr Gln Ala Cys  
 160 165  
 CCA AAG ACA TCC TTT GAG CCA ATT CCC ATA CAT TAT TGT 544  
 Pro Lys Thr Ser Phe Glu Pro Ile Pro Ile His Tyr Cys  
 170 175 180  
 10 GCC CCG GCT GGT TTT GCG ATT CTA AAG TGT AAT GAT AAG 583  
 Ala Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asp Lys  
 185 190  
 AAG TTC AAT GGA ACA GGA CCA TGT AAA AAT GTC AGC ACA 622  
 Lys Phe Asn Gly Thr Gly Pro Cys Lys Asn Val Ser Thr  
 195 200 205  
 GTA CAA TGT ACA CAT GGA ATT AGG CCA GCA GTA TCA ACT 661  
 Val Gln Cys Thr His Gly Ile Arg Pro Val Val Ser Thr  
 210 215 220  
 CAA CTG TTG TTA AAT GGC ACT CTA GCA GAA GAG GTA 700  
 20 Gln Leu Leu Asn Gly Ser Leu Ala Glu Glu Glu Val  
 225 230  
 GTA ATC AGA TCT GCC AAT TTC ACA GAC AAT GCT AAA ACC 739  
 Val Ile Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr  
 235 240 245  
 25 ATA ATA GTA CAT CTA AAT GAA ACT GTA AAA ATT AAT TGT 778  
 Ile Ile Val His Leu Asn Glu Thr Val Lys Ile Asn Cys  
 250 255  
 ACA AGA CTT GGC AAC AAT ACA AGA AAA AGT ATA AAT ATA 817  
 Thr Arg Leu Gly Asn Asn Thr Arg Lys Ser Ile Asn Ile  
 30 260 265 270  
 GGA CCA GGG AGA GTA CTC TAT GCA ACA GGA GAA ATA ATA 856  
 Gly Pro Gly Arg Val Leu Tyr Ala Thr Gly Glu Ile Ile  
 275 280 285  
 GGA GAC ATA AGA CAA GCA CAT TGT AAC ATT AGT AGA GCA 895  
 35 Gly Asp Ile Arg Gln Ala His Cys Asn Ile Ser Arg Ala  
 290 295  
 CAA TGG AAT AAG ACT TTA GAA AAG GTA GTT GAC AAA TTA 934  
 Gln Trp Asn Lys Thr Leu Glu Lys Val Val Asp Lys Leu  
 300 305 310  
 40 AGA AAA CAA TTT GGG GAT AAT ACA ACA ATA GCT TTT AAT 973  
 Arg Lys Gln Phe Gly Asp Asn Thr Ile Ala Phe Asn  
 315 320  
 CGA TCC TCA GGA GGG GAC CCA GAA ATT GTA ATG CAC ACT 1012  
 Arg Ser Ser Gly Gly Asp Pro Glu Ile Val Met His Thr  
 45 325 330 335  
 TTT AAT TGT GGA GGG GAA TTT TTC TAC TGT AAT ACA ACA 1051  
 Phe Asn Cys Gly Gly Glu Phe Phe Tyr Cys Asn Thr Thr  
 340 345 350  
 CAA CTG TTT AAT AGT ACT TGG AAT AAT ACT TGG AAG GAT 1090  
 50 Gln Leu Phe Asn Ser Thr Trp Asn Asn Thr Trp Lys Asp  
 355 360  
 CCT AAC AGG AGT GAC AAT ATC ACA CTC CCA TGC AGA ATA 1129  
 Pro Asn Arg Ser Asp Asn Ile Thr Leu Pro Cys Arg Ile  
 365 370 375  
 55 AAA CAA ATT ATA AAC ATG TGG CAG GAA GTA GGA AAA GCA 1168  
 Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys Ala  
 380 385  
 ATG TAC GCC CCT CCC ATC AGA GGG GAA ATT AGA TGT TCA 1207  
 Met Tyr Ala Pro Pro Ile Arg Gly Glu Ile Arg Cys Ser  
 60 390 395 400  
 TCA AAT ATC ACA GGG CTG CTA CTA ACA AGA GAT GGT GGT 1246  
 Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly  
 405 410 415

AAT GAC GAT GGT AAT GAC ACG ACC ACA AAC AGG ACC GAG 1285  
 Asn Asp Asp Gly Asn Asp Thr Thr Thr Asn Arg Thr Glu  
 420 425  
 ATC TTC AGA CCT GGA GGA GGA GAT ATG AGG GAC AAT TGG 1324  
 5 Ile Phe Arg Pro Gly Gly Asp Met Arg Asp Asn Trp  
 430 435 440  
 AGA AGT GAA TTA TAT AGA TAT AAA GCA GCA AAA ATT GAA 1363  
 Arg Ser Glu Leu Tyr Arg Tyr Lys Val Val Lys Ile Glu  
 445 450  
 10 CCA TTA GGA ATA GCA CCC ACC AGG GCA AAG AGA AGA GTG 1402  
 Pro Leu Gly Ile Ala Pro Thr Arg Ala Lys Arg Arg Val  
 455 460 465  
 GTG CAG AGA GAA AAA AGA GCA GCA GGA CTA GGA GCT TTG 1441  
 Val Gln Arg Glu Lys Arg Ala Val Gly Leu Gly Ala Leu  
 15 470 475 480  
 TTC CTT GGG T TCTTAGGAG CATAAAGCTT CTAGA 1475  
 Phe Leu Gly  
 483

**20 CLONE C10.7**  
 G GTA CCT GTG TGG AAA GAA GCA AAC ACA ACT CTA TTT 37  
 Val Pro Val Trp Lys Glu Ala Asn Thr Thr Leu Phe  
 1 5 10  
 TGT GCA TCA GAT GCT AAA GCA TAT GAT AGA GAA GTA CAT 76  
 25 Cys Ala Ser Asp Ala Lys Ala Tyr Asp Arg Glu Val His  
 15 20 25  
 AAT GTT TGG GCA ACA CAT GCC TGT GTA CCC ACA GAC CCC 115  
 Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro  
 30 35  
 AAC CCA CAA GAA ATA GTA TTG GGA AAT GTG ACA GAA AAT 154  
 Asn Pro Gln Glu Ile Val Leu Gly Asn Val Thr Glu Asn  
 40 45 50  
 TTT AAC ATG TGG AAA AAT AAC ATG GTA GAA CAA ATG CAT 193  
 Phe Asn Met Trp Lys Asn Asn Met Val Glu Gln Met His  
 35 55 60  
 GAG GAT ATA ATC AAT TTA TGG GAT CAA AGC TTA AAG CCA 232  
 Glu Asp Ile Ile Asn Leu Trp Asp Gln Ser Leu Lys Pro  
 65 70 75  
 TGT GTA AAG TTA ACT CCA CTC TGT GTT ACT TTA AAG TGC 271  
 40 Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu Lys Cys  
 80 85 90  
 AAG GAT CTG GAG AGC AAT ACT ACC TAT AAT AGC ACT ATT 310  
 Lys Asp Leu Glu Arg Asn Thr Thr Tyr Asn Ser Thr Ile  
 95 100  
 45 ACC AAT AAT AGT AGT TTG GAG GGA CTA AGA GAA CAA ATG 349  
 Thr Asn Asn Ser Ser Leu Glu Gly Leu Arg Glu Gln Met  
 105 110 115  
 ACA AAC TGC TCT TTC AAC ATC ACC ACA AGT ATA AGA GAT 388  
 Thr Asn Cys Ser Phe Asn Ile Thr Thr Ser Ile Arg Asp  
 50 120 125  
 AAG GTG CAG AAA GAA TAT GCA CTT TTG TAT AAA CTT GAT 427  
 Lys Val Gln Lys Glu Tyr Ala Leu Leu Tyr Lys Leu Asp  
 130 135 140  
 GTA GTA CCA ATA GAA GAA GAT GAC AAT ACT AGC TAT AGA 466  
 55 Val Val Pro Ile Glu Glu Asp Asp Asn Thr Ser Tyr Arg  
 145 150 155  
 TTG ATA AGT TGT AAC ACC TCA GTC ATT ACA CAG GCT TGT 505  
 Leu Ile Ser Cys Asn Thr Ser Val Ile Thr Gln Ala Cys  
 160 165  
 60 CCA AAG ACA TCC TTT GAG CCA ATT CCC ATA CAT TAT TGT 544  
 Pro Lys Thr Ser Phe Glu Pro Ile Pro Ile His Tyr Cys  
 170 175 180  
 GCC CCG GCT GGT TTT GCG ATT CTA AAG TGT AAT GAT AAG 583  
 Ala Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asp Lys  
 65 185 190

1 AAG TTC AAT GGA ACA GGA CCA TGT AAA AAT GTC AGC ACA 622  
 Lys Phe Asn Gly Thr Gly Pro Cys Lys Asn Val Ser Thr  
 195 200 205  
 5 CTA CAA TGT ACA CAT GGA ATT AGG CCA GTC GTC TCA ACT 661  
 Val Gln Cys Thr His Gly Ile Arg Pro Val Val Ser Thr  
 210 215 220  
 CAA CTG TTG TTA AAT GGC AGT CTA GCA GAA GAG GTC 700  
 Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu Glu Val  
 225 230  
 10 GTC ATC AGA TCT GCC AAT TTC ACA GAC AAT GCT AAA ACC 739  
 Val Ile Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr  
 235 240 245  
 ATA ATA GTA CAT CTA AAT GAA ACT GTC AAA ATT AAT TGT 778  
 Ile Ile Val His Leu Asn Glu Thr Val Val Lys Ile Asn Cys  
 15 250 255  
 ACA AGA CTT GGC AAC AAT ACA AGA AAA AGT ATA AAT ATA 817  
 Thr Arg Leu Gly Asn Asn Thr Arg Lys Ser Ile Asn Ile  
 260 265 270  
 GGA CCA GGG AGA GTC CTC TAT GCA ACA GGA GAA ATA ATA 856  
 20 Gly Pro Gly Arg Val Leu Tyr Ala Thr Gly Glu Ile Ile  
 275 280 285  
 GGA GAC ATA AGA CAA GCA CAT TGT AAC ATT AGT AGA GCA 895  
 Gly Asp Ile Arg Gln Ala His Cys Asn Ile Ser Arg Ala  
 290 295  
 25 CAA TGG AAT AAG ACT TTA GAA AAG GTC GTT GAC AAG TTA 934  
 Gln Trp Asn Lys Thr Leu Glu Lys Val Val Asp Lys Leu  
 300 305 310  
 AGA AAA CAA TTT GGG GAT AAT ACA ACA ATA GCT TTT AAT 973  
 Arg Lys Gln Phe Gly Asp Asn Thr Thr Ile Ala Phe Asn  
 315 320  
 CGA TCC TCA GGA GGG GAC CCA GAA ATT GTC ATG CAC ACT 1012  
 Arg Ser Ser Gly Gly Asp Pro Glu Ile Val Met His Thr  
 325 330 335  
 TTT AAT TGT GGA GGG GAA TTT TTC TAC TGT AAT ACA ACA 1051  
 35 Phe Asn Cys Gly Gly Glu Phe Phe Tyr Cys Asn Thr Thr  
 340 345 350  
 CAA CTG TTT AAT AGT ACT TGG AAT AAT ACT TGG AAG GAT 1090  
 Gln Leu Phe Asn Ser Thr Trp Asn Asn Thr Trp Lys Asp  
 355 360  
 40 CCT AAC AGG AGT GAC AAT ATC ACA CTC CCA TGC AGA ATA 1129  
 Pro Asn Arg Ser Asp Asn Ile Thr Leu Pro Cys Arg Ile  
 365 370 375  
 AAA CAA ATT ATA AAC ATG TGG CAG GAA GTC GGA AAA CCA 1168  
 Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys Ala  
 380 385  
 45 ATG TAC GCC CCT CCC ATC AGA GGG GAA ATT AGA TGT TCA 1207  
 Met Tyr Ala Pro Pro Ile Arg Gly Glu Ile Arg Cys Ser  
 390 395 400  
 TCA AAT ATC ACA GGG CTG CTA ACA AGA GAT GGT GGT 1246  
 50 Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly  
 405 410 415  
 AAT GAC GAT GGT AAT GAC ACG ACC ACA AAC AGG ACC GAG 1285  
 Asn Asp Asp Gly Asn Asp Thr Thr Asn Arg Thr Glu  
 420 425  
 55 ATC TTC AGA CCT GGA GGA GAT ATG AGG GAC AAT TGG 1324  
 Ile Phe Arg Pro Gly Gly Asp Met Arg Asp Asn Trp  
 430 435 440  
 AGA AGT GAA TTA TAT AGA TAT AAA GTC GTC AAA ATT GAA 1363  
 Arg Ser Glu Leu Tyr Arg Tyr Lys Val Val Lys Ile Glu  
 60 445 450  
 CCA TTA GGA ATA GCA CCC ACC AGG GCA AAG AGA AGA GTG 1402  
 Pro Leu Gly Ile Ala Pro Thr Arg Ala Lys Arg Arg Val  
 455 460 465

5' GTG CAG AGA GAA AAA ACA GCA GTA GGA CTA GGA GCT TTG 1441  
 Val Gln Arg Glu Lys Arg Ala Val Gly Leu Gly Ala Leu  
 470 475 480  
 TTC CTT GGG TTC TTG GGA GCA TAA AGC TTC TAG A 1475  
 5' Phe Leu Gly Phe Leu Gly Ala Xaa Ser Phe Xaa  
 485 490 491

CLONE C17.1

10' CTC GAG GTA CCT GTG TGG AAA GAA GCA ACC ACC ACT 36  
 Leu Glu Val Pro Val Trp Lys Glu Ala Thr Thr Thr  
 1 5 10  
 CTA TTT TGT GCA TCA GAT GCT AAA GCA TAT GAT TCA GAG 75  
 Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Ser Glu  
 15 20 25  
 15' GCA CAT AAT GTT TGG GCC ACA CAT GCC TGT GTA CCC ACA 114  
 Ala His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr  
 30 35  
 GAC CCC AAC CCA CAA GAA GTC GAA TTG GAA AAT GTG ACA 153  
 Asp Pro Asn Pro Gln Glu Val Glu Leu Glu Asn Val Thr  
 20 40 50  
 GAA AAT TTT AAC ATG TGG AAA AAT AAC ATG GTA GAA CAG 192  
 Glu Asn Phe Asn Met Trp Lys Asn Asn Met Val Glu Gin  
 55 60  
 ATG CAT GGG GAT ATA ATT ACT TTA TGG GAT CAA AGC CTA 231  
 25' Met His Gly Asp Ile Ile Ser Leu Trp Asp Gln Ser Leu  
 65 70 75  
 AAG CCA TGT GTA AAA TTA ACC CCA CTC TGT GTT ACG TTA 270  
 Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu  
 80 85 90  
 30' AAT TGC ACT GAC CCA AAT GTT ACT AAT AGC GAG AGA AGC 309  
 Asn Cys Thr Asp Pro Asn Val Thr Asn Ser Glu Arg Thr  
 95 100  
 ATA GAG GGG GGA GAA ATA AAA AAT TGC TCT TTC AAT ATC 348  
 Ile Glu Gly Gly Glu Ile Lys Asn Cys Ser Phe Asn Ile  
 35 105 110 115  
 ACC ACA AAC ATA AGA GAT AGG TTT CAG AAA GAA TAT GCA 387  
 Thr Thr Asn Ile Arg Asp Arg Phe Gln Lys Glu Tyr Ala  
 120 125  
 CTT TTT TAT AAA CTT GAT GTA ATA CCA TTA GGT AAT GAT 426  
 40' Leu Phe Tyr Lys Leu Asp Val Ile Pro Leu Gly Asn Asp  
 130 135 140  
 AAT ACT AGC TAT AGG TTG ATA AGT TGT AAC ACC TCA GTC 465  
 Asn Thr Ser Tyr Arg Leu Ile Ser Cys Asn Thr Ser Val  
 145 150 155  
 45' ATT ACA CAG GCC TGT CCA AAG GTA TCC TTT GAG CCA ATT 504  
 Ile Thr Gln Ala Cys Pro Lys Val Ser Phe Glu Pro Ile  
 160 165  
 CCC ATA CAT TAT TGT GCC CCG GCT GGT TTT GCG ATT CTA 543  
 Pro Ile His Tyr Cys Ala Pro Ala Gly Phe Ala Ile Leu  
 50 170 175 180  
 AAG TGT AAA GAT AAG AAG TTC AAT GGA ACA GGA CCA TGT 582  
 Lys Cys Lys Asp Lys Phe Asn Gly Thr Gly Pro Cys  
 185 190  
 ACA AAT GTC AGC ACA GTA CAA TGT ACA CAT GGA ATT AAG 621  
 55' Thr Asn Val Ser Thr Val Gln Cys Thr His Gly Ile Lys  
 195 200 205  
 CCA GTA GTA TCA ACT CAA CTG TTG TTA AAT GGC AGT CTA 660  
 Pro Val Val Ser Thr Gln Leu Leu Leu Asn Gly Ser Leu  
 210 215 220  
 60' GCA GAA GAA GAC ATA GTA ATT AGA TCC GCC AAT CTC ACA 699  
 Ala Glu Glu Asp Ile Val Ile Arg Ser Ala Asn Leu Thr  
 225 230  
 GAC AAT GCT AAA AAC ATA ATA GTA CAG CTG AAT GAA TCT 738  
 Asp Asn Ala Lys Asn Ile Ile Val Gln Leu Asn Glu Ser  
 65 235 240 245

GTA ACA ATG AAT TGT ACA AGA CCC AAC AAC AAT ACA ATG 777  
 Val Thr Met Asn Cys Thr Arg Pro Asn Asn Asn Thr Met  
 250 255  
 5 AAA AGT ATA CAT ATA GGA CCA GGC AGA GCA TTT TAT GCA 816  
 Lys Ser Ile His Ile Gly Pro Gly Arg Ala Phe Tyr Ala  
 260 265 270  
 ACA GGA AAC ATA ATA GGA GAT ATA AGA CAA GCA CAT TGT 855  
 Thr Gly Asn Ile Ile Gly Asp Ile Arg Gln Ala His Cys  
 275 280 285  
 10 AAC ATT AGT GGA ACA AAA TGG AAT GAC ACT TTG AAA AAG 894  
 Asn Ile Ser Gly Thr Lys Trp Asn Asp Thr Leu Lys Lys  
 290 295  
 ATA GCT ATA AAA TTA AGA GAA CAA TTT AAT AAG ACA ATA 933  
 Ile Ala Ile Lys Leu Arg Glu Gln Phe Asn Lys Thr Ile  
 300 305 310  
 15 GTC TTT AAT CAA TCC TCA GGA GGG GAC CCA GAA ATT GCA 972  
 Val Phe Asn Gln Ser Ser Gly Gly Asp Pro Glu Ile Ala  
 315 320  
 20 ACG CTC AGT TTT AAT TGT GGA GGG GAA TTT TTC TAC TGT 1011  
 Thr Leu Ser Phe Asn Cys Gly Gly Glu Phe Phe Tyr Cys  
 325 330 335  
 AAT TCA ACA CAA CTG TTT AAT AGT ACT TGG AAT AGT ACT 1050  
 Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp Asn Ser Thr  
 340 345 350  
 25 GGG TCA AAT AAC ACT AAA GGA AAT GAC ACA ATC ACA CTC 1089  
 Gly Ser Asn Asn Thr Lys Gly Asn Asp Thr Ile Thr Leu  
 355 360  
 CCA TGC AGA ATA AGA CAA ATT ATA AAC ATG TGG CAG AAA 1128  
 Pro Cys Arg Ile Arg Gln Ile Ile Asn Met Trp Gln Lys  
 365 370 375  
 30 ATA GGA AAA GCA ATG TAT GCC CCT CCC ATC AAA GGG CAA 1167  
 Ile Gly Lys Ala Met Tyr Ala Pro Pro Ile Lys Gly Gln  
 380 385  
 35 ATT AGA TGT TCA TCA AAT ATT ACA GGG CTA ATA TTA ACA 1206  
 Ile Arg Cys Ser Ser Asn Ile Thr Gly Leu Ile Leu Thr  
 390 395 400  
 AGA GAT GGT GGT AAC AAC AAC ATG ACC AAG ACC ACC GAG 1245  
 Arg Asp Gly Gly Asn Asn Asn Met Ser Lys Thr Thr Glu  
 405 410 415  
 40 ACC TTC AGA CCT GGA GGA GAT ATG AGC GAC AAT TGG 1284  
 Thr Phe Arg Pro Gly Gly Asp Met Arg Asp Asn Trp  
 420 425  
 AGA AGT CAA TTA TAT AAA TAT AAA GTC GTC AAA ATT GAA 1323  
 Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile Glu  
 430 435 440  
 45 CCA TTA GGA GTA GCA CCC ACC AGG GCA AAG AGA AGA GTG 1362  
 Pro Leu Gly Val Ala Pro Thr Arg Ala Lys Arg Arg Val  
 445 450  
 GTG CAG AGA GAA AAA AGA GCA GTG GGA ATA GGA GCT GTG 1401  
 50 Val Gln Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Val  
 455 460 465  
 TTC CTT GGG TTC TTG GGA GCA TAA AGC TTC TAG A 1435  
 Phe Leu Gly Phe Leu Gly Ala Xaa Ser Phe Xaa  
 470 475 478

55

CLONE C17.3

CTC GAG GTA CCT GTG TGG AAA GAA GCA ACC ACC ACT 36  
 Leu Glu Val Pro Val Trp Lys Glu Ala Thr Thr Thr  
 1 5 10  
 60 CTA TTT TGT GCA TCA GAT GCT AAA GCA TAT GAT TCA GAG 75  
 Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Ser Glu  
 15 20 25  
 GCA CAT AAT GTT TGG GCC ACA CAT GCC TGT GTA CCC ACA 114  
 Ala His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr  
 65 30 35



GTC TTT AAT CAA TCC TCA GGA GGG GAC CCA GAA ATT GCA 972  
 Val Phe Asn Gln Ser Ser Gly Gly Asp Pro Glu Ile Ala  
 315 320  
 ACG CTC AGT TTT AAT TGT GGA GGG GAA TTT TTC TAC TGT 1011  
 5 Thr Leu Ser Phe Asn Cys Gly Gly Glu Phe Phe Tyr Cys  
 325 330 335  
 AAT TCA ACA CAA CTG TTT AAT ACT ACT TGG AAT AGT ACT 1050  
 Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp Asn Ser Thr  
 340 345 350  
 10. GGG TCA AAT AAC ACT AAA GGA AAT GAC ACA ATC ACA CTC 1089  
 Gly Ser Asn Asn Thr Lys Gly Asn Asp Thr Ile Thr Leu  
 355 360  
 CCA TGC AGA ATA AGA CAA ATT ATA AAC ATG TGG CAG AAA 1128  
 Pro Cys Arg Ile Arg Gln Ile Ile Asn Met Trp Gln Lys  
 15. 365 370 375  
 ATA GGA AAA GCA ATG TAT GCC CCT CCC ATC AAA GGG CAA 1167  
 Ile Gly Lys Ala Met Tyr Ala Pro Pro Ile Lys Gly Gln  
 380 385  
 ATT AGA TGT TCA TCA AAT ATT ACA GGG CTA ATA TTA ACA 1206  
 20. Ile Arg Cys Ser Ser Asn Ile Thr Gly Leu Ile Leu Thr  
 390 395 400  
 AGA GAT GGT GGT AAC AAC AAC ATG AGC AAG ACC ACC GAG 1245  
 Arg Asp Gly Gly Asn Asn Asn Met Ser Lys Thr Thr Glu  
 405 410 415  
 25. ACC TTC AGA CCT GGA GGA CGA GAT ATG AGG GAC AAT TGG 1284  
 Thr Phe Arg Pro Gly Gly Asp Met Arg Asp Asn Trp  
 420 425  
 AGA AGT GAA TTA TAT AAA TAT AAA GTC GTC AAA ATT GAA 1323  
 Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile Glu  
 30. 430 435 440  
 CCA TTA GGA GTC GCA CCC ACC AGG GCA AAG AGA AGA GTG 1362  
 Pro Leu Gly Val Ala Pro Thr Arg Ala Lys Arg Arg Val  
 445 450  
 GTG CAG AGA GAA AAA AGA GCA GTG GGA ATA GGA GCT GTG 1401  
 35. Val Gln Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Val  
 455 460 465  
 TTC CTT GGG TTC TTG GGA GCA TAA AGC TTC TAG A 1435  
 Phe Leu Gly Phe Leu Gly Ala Xaa Ser Phe Xaa  
 470 475 478

40.

In addition to the listing in Table 1, Figure 3 shows  
 the alignment of the amino acid sequences of the clones  
 of each of the seven isolates. Corresponding residues  
 from various clones are in boxes. In the figure, the  
 45. amino acid sequences are aligned against MN-rgp120

(SEQ. ID. NO. 29).

In one embodiment, a gp120 polypeptide of this  
 invention has the same amino acid sequence as the  
 sequence of one of the breakthrough isolates. In  
 50. another embodiment, the amino acid sequence is  
 truncated, as described in detail hereinafter. In  
 another embodiment, a gp120 polypeptide sequence of  
 this invention contains a substitution, insertion, or

deletion (alteration) of one or more amino acids in the sequence of a breakthrough isolate. Usually, with the exception of amino acids that are not present in a truncated amino acid sequence and eliminate an epitope,

5 a gp120 polypeptide of this invention will include alterations in the amino acid sequence of a breakthrough isolate that do not alter the polypeptide's ability to induce the same neutralizing antibodies as the amino acid sequence of the isolate.

10 In general, substitutions in the amino acid sequence of a gp120 polypeptide of this invention are conservative substitutions, particularly for amino acid residues in the V2, V3, and C4 domains of gp120, which domains contain neutralizing epitopes. However, non-conservative substitutions, particularly in domains

15 that do not contain neutralizing epitopes are contemplated.

20 Conservative substitutions replace an amino acid with an amino acid of similar size and character. For example, a hydrophobic residue or hydrophilic residue is replaced with another hydrophobic residue or hydrophilic residue, respectively. Amino acids can be divided into the following groups: positively charged residues (K, R and H); negatively charged residues

25 (D and E); amides (N and Q); aromatics (F, Y, and W); hydrophobics (P, G, A, V, L, I, and M); and uncharged residues (S and T). Usually, residues within a group are replaced with another member of the group.

30 In one embodiment, critical amino acid residues in the V2, V3, and C4 domains of gp120 are identical to the corresponding residues in a breakthrough isolate sequence. Critical amino acid residues in the V2, V3, and C4 domains of gp120 are described in the experimental section. In another embodiment, all amino

acid residues in the V2, V3, and C4 domains of gp120 are identical to corresponding residues in a breakthrough isolate sequence.

5 5 Oligonucleotide Encoding gp120 from Breakthrough Isolates

The present invention also provides novel oligonucleotides encoding gp120 from the breakthrough isolates which can be used to express gp120. An 10 oligonucleotide of this invention encodes a polypeptide of this invention. The oligonucleotide can be DNA or RNA, usually DNA. Although numerous nucleotide sequences can encode the same amino acid sequence due to the degeneracy of the genetic code, conveniently, 15 the oligonucleotides of this invention include a nucleotide sequence of a breakthrough isolate as illustrated in Table 1 (Sequence ID Nos. 2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, 24, 26, and 28). Usually, an oligonucleotide of this invention is less than about 20 5 kilobases (kb), preferably less than about 3 kb.

To express the encoded amino acid sequence, the oligonucleotide can be inserted into a transcription unit. The transcription unit can be inserted into a plasmid for production of cell lines, inserted into a 25 virus (e.g.; vaccinia) or can be used directly as a DNA vaccine. Suitable transcription units for production of vaccine proteins are well known. A preferred expression vector, designated psvI6B5, is illustrated in Sequence ID No. 32. The vector includes an HSV-1 30 gD1 signal sequence joined to a linker sequence. The gp120 nucleotide sequence to be expressed starts with the Kpn I site of the gene. Since all gp120 or gp160 sequences contain this site, any gp120 nucleotide sequence can be analogously inserted into the vector 35 and expressed. The vector ends with a polyA tail from SV40.

In addition to being useful to express a polypeptide sequence of this invention, the oligonucleotides of this invention can also be used in diagnostics to detect HIV isolates. For example, the 5 oligonucleotide or a portion thereof encoding a neutralizing epitope can be used in branched chain DNA diagnostics or as a probe in in situ hybridization studies.

10 Vaccine preparation

A gp120 polypeptide of this invention from a selected breakthrough isolate(s) in a suitable carrier is used to make a subunit vaccine. The polypeptide can be used alone, but is generally administered in a 15 multivalent subunit vaccine that includes gp120 MN. In addition to one or more gp120 polypeptides of this invention, the vaccine generally includes the MN polypeptide (hereinafter, MN-rgp120). The vaccine usually includes about 3 to about 5 different gp120 20 polypeptides, but 30 or more different gp120 polypeptides can be used.

Preparation of gp120 polypeptides for use in a vaccine is well known and is described hereinafter. With the exception of the use of the selected HIV 25 isolate, the gp120 subunit vaccine prepared in the method does not differ from gp120 subunit vaccines of the prior art.

As with prior art gp120 subunit vaccines, gp120 at the desired degree of purity and at a sufficient 30 concentration to induce antibody formation is mixed with a physiologically acceptable carrier. A physiologically acceptable carrier is nontoxic to a recipient at the dosage and concentration employed in the vaccine. Generally, the vaccine is formulated for 35 injection, usually intramuscular or subcutaneous injection. Suitable carriers for injection include

sterile water, but preferably are physiologic salt solutions, such as normal saline or buffered salt solutions such as phosphate-buffered saline or ringer's lactate. The vaccine generally contains an adjuvant.

5      Useful adjuvants include QS21 (Quillaja saponaria, commercially available from Cambridge Biotech, Worcester, MA), which stimulates cytotoxic T-cells, and alum (aluminum hydroxide adjuvant). Formulations with different adjuvants which enhance cellular or local

10     immunity can also be used. In particular, immunopotentiators such as cytokines can be included in the vaccine. Examples of suitable immunopotentiating cytokines include interleukins, such as interleukin-2 (IL-2) and interleukin-12 (IL-12), and tumor necrosis

15     factor-alpha (TNF- $\alpha$ ).

Additional excipients that can be present in the vaccine include low molecular weight polypeptides (less than about 10 residues), proteins, amino acids, carbohydrates including glucose or dextrans, chelating agents such as EDTA, and other excipients that stabilize the protein or inhibit growth of microorganisms.

The vaccine can also contain other HIV proteins. In particular, gp41 or the extracellular portion of gp41 or HIV-1 core proteins such as P24, P17, and P55 can be present in the vaccine. Although the amino acid sequence of gp41 is more conserved than that of gp120, gp41 contains neutralizing epitopes. Preferably, any gp41 present in the vaccine is from an HIV isolate present in the vaccine. gp160 from an isolate used in the vaccine can replace gp120 in the vaccine or be used together with gp120 from the isolate. Alternatively, gp160 from a different isolate than those in the vaccine can additionally be present in the vaccine.

35     Vaccines according to the invention can also contain one or more soluble gp120 polypeptide

sequences, or fragments thereof, in combination with an engineered virus specifically designed to express proteins that induce a cytotoxic T-cell response. Suitable engineered viruses are derived from, for 5 example, Canary Pox virus, vaccinia viruses, attenuated human herpes viruses (such as, e.g., herpes simplex viruses), and Varicella Zoster. Exemplary engineered viruses are modified to express any HIV protein capable of inducing a cytotoxic T-cell response, such as those 10 described above. Typically, immunization with the gp120/engineered virus vaccine is followed by administration of one or more doses of the gp120 polypeptide sequence(s) to boost the immune response. If desired, viruses can be engineered to express one or 15 more gp120 polypeptide sequences of the invention, or fragments thereof, and used in vaccines with or without soluble gp120 polypeptide sequences.

Vaccine formulations generally include a total of about 300 to 600  $\mu$ g of gp120, conveniently in about 20 1.0 ml of carrier. Preferred formulations include use of twice the weight of a gp120 polypeptide in twice as 600  $\mu$ g alum. However, formulations having smaller amounts (e.g.; 50  $\mu$ g per dose) are also used, generally with alum or other adjuvants. The amount of gp120 for 25 any isolate present in the vaccine will vary depending on the immunogenicity of the gp120. For example, gp120 from some strains of HIV may be less immunogenic than gp120 from the MN strain (Sequence ID No. 29). If two strains having different immunogenicity are used in 30 combination, empirical titration of the amount of each virus would be performed to determine the percent of the gp120 of each strain in the vaccine. For isolates having similar immunogenicity, approximately equal amounts of each isolate's gp120 would be present in the 35 vaccine. For example, in a preferred embodiment, the vaccine includes gp120 from the MN and a strain of this

invention at concentrations of about 300  $\mu$ g per strain in about 1.0 ml of carrier. When the vaccine includes gp120 from about 30 isolates, about 10 to about 50  $\mu$ g can be used. Methods of determining the relative 5 amount of an immunogenic protein in multivalent vaccines are well known and have been used, for example, to determine relative proportions of various isolates in multivalent polio vaccines.

The vaccines of this invention are administered in 10 the same manner as prior art HIV gp120 subunit vaccines. In particular, the vaccines are generally administered at 0, 1, and at 6, 8 or 12 months, depending on the protocol. A preferred protocol includes administration at 0, 1, 6, and 12 months. 15 Following the immunization procedure, annual or bi-annual boosts can be administered. However, during the immunization process and thereafter, neutralizing antibody levels can be assayed and the protocol adjusted accordingly.

20 The vaccine is administered to uninfected individuals. In addition, the vaccine can be administered to seropositive individuals to augment immune response to the virus, as with prior art HIV vaccines. It is also contemplated that DNA encoding 25 the strains of gp120 for the vaccine can be administered in a suitable vehicle for expression in the host. In this way, gp120 can be produced in the infected host, eliminating the need for repeated immunizations. Preparation of gp120 expression 30 vehicles is described hereinafter.

35 Although the gp120 isolates described herein can be used as a vaccine as described above, the amino acid sequences can also be used alone or in combinations in the same type of formulation for use as an immunogen, to induce antibodies that recognize the isolate(s) present in the immunogen. Immunogens are formulated in

the same manner as vaccines and can include the same excipients, etc. Antibodies induced by the immunogens can be used in a diagnostic to detect the HIV strain in the immunogen or to affinity purify the strain.

5

#### gp120 Polypeptide Sequences and Chemokine Receptors

While CD4 is the primary cellular receptor for HIV-1, it is not sufficient for entry of HIV-1 into cells. Co-receptors required in conjunction with CD4 10 have been identified. These co-receptors are members of the chemokine receptor family of seven-transmembrane G-protein coupled receptors. The chemokine superfamily is subdivided into two groups based on the amino terminal cysteine spacing. The CXC chemokines are 15 primarily involved in neutrophil-mediated inflammation, and the CC chemokines tend to be involved in chronic inflammation. At least five CC chemokine receptors, designated CC-CKR1-5 (also known in the art as CCR1-5), and at least four CXC chemokine receptors, designated 20 CXC-CKR1-4 (also known as CXCR-1-4), have been identified.

CXC-CKR-4 (CXCR-4), which has also been called the alpha-chemokine receptor fusin, serves as an entry cofactor for T-cell-tropic HIV-1 strains. CC-CKR-5 25 (CC-R5), which has been called beta-chemokine receptor, together with its related family members, such as CC-CKR-2b and CC-CKR3, serve as entry cofactors for macrophage-tropic HIV-1 strains. T-cell-tropic strains can infect primary T-cells and T-cell lines, but not 30 macrophages, whereas macrophage-tropic strains can infect macrophages and primary T-cells, but not T-cell lines. T-cell- and macrophage-tropic strains are discussed more fully in Deng et. al., Nature 381:661-666 (1996), which is hereby incorporated by reference 35 in its entirety. Examples of T-cell-tropic strains include laboratory isolates, such as IIIB and MN.

Macrophage-tropic strains include primary isolates, including but not limited to A244, GNE6, GNE8, and breakthrough viruses from vaccinees immunized with gp120-based vaccines. Dual-tropic strains can use both 5 types of co-receptors, entering cells via CXC-CKR-4 or via one or more CC-CKR family members, preferably CC-CKR-5, CC-CKR-2b, or CC-CKR-3. While the present invention is not intended to be bound or limited by any one theory, the entry of T-cell tropic and macrophage-tropic HIV-1 strains is believed to provide a unifying 10 explanation of the differences in cell tropism between viral strains, the resistance to HIV-1 infection by many CD4-transfected nonprimate cells, and the HIV-1 infection resistance of a portion of the human 15 population.

Accordingly, in one embodiment is a vaccine containing (1) a first gp120 polypeptide sequence, or fragment thereof, from a macrophage-tropic HIV-1 strain and/or a second gp120 polypeptide sequence, or fragment 20 thereof, from a T-cell tropic strain, in combination with (2) a breakthrough isolate HIV gp120 polypeptide sequence, or fragment thereof, from a vaccinee vaccinated with the first and/or second HIV gp120 25 polypeptide sequence. Preferably, the vaccine includes at least two gp120 polypeptide sequences that bind to different chemokine receptors. In one embodiment, the vaccine includes first and second gp120 polypeptide sequences that bind to different chemokine receptors. In addition, the breakthrough isolate gp120 polypeptide 30 sequence can bind to a different chemokine receptor than the chemokine receptor(s) bound by either or both of the first and second gp120 polypeptide sequence(s).

A preferred T-cell tropic strain is a laboratory isolate, most preferably MN. Preferred macrophage-tropic viruses for use in the invention are GNE6 and 35 GNE8, which are representative of the breakthrough

5 viruses disclosed herein and differ from MN in that their gp120s induce the formation of antibodies that recognize the gp120 sequences (e.g., the V3 domain) involved in binding to CC chemokine receptors, such as CXC-CKR-5.

In one embodiment, HIV infection is prevented by administering one or more chemokine receptor-binding gp120 polypeptide sequences, or fragment(s) thereof containing appropriate chemokine receptor-binding domains, in a vaccine, such as those described above. Preferably, the vaccine also includes one or more CD4-binding gp120 polypeptide sequences or appropriate fragments thereof. Such vaccines induce anti-HIV antibodies that inhibit viral gp120-chemokine receptor or -CD4 binding. In addition, such gp120 polypeptides can directly inhibit HIV infection by binding to one or more co-receptors for HIV infection, such as CD4 or a chemokine receptor, thus providing a prophylactic or therapeutic effect in treating HIV infection. Preferably, gp120 polypeptide sequences useful in this regard contain the T-cell binding (TCB) domain.

Various uses of chemokine receptor-binding gp120 polypeptides are discussed below with regard to the CC chemokine receptor family. However, those skilled in the art recognize that this discussion applies equally to CXC chemokine receptors that act as cofactors in HIV infection.

The gp120 polypeptides can be used as a composition containing one or more gp120 polypeptides, as described for use as a vaccine or immunogen. The composition can be administered, prophylactically or therapeutically, to a patient at risk of infection or in need of such treatment using the dosages and routes and means of administration described herein. However, chronic administration may be preferred and dosages can be adjusted accordingly. It is noted that *in vivo*

administration can also induce antibodies that bind viral gp120, further inhibiting virus binding to CC-CKR.

The gp120 polypeptides can also be used in 5 screening assays to identify antagonists of CC-CKR.

For example, candidate antagonists can be screened for inhibition of binding of gp120 to a CC-CKR CC-CKR receptor that is isolated and attached to a surface (e.g., plastic dish) or recombinantly or naturally 10 expressed on the surface of a cell. Antagonists can either bind gp120 or bind receptor. Preferred candidate antagonists include gp120 compounds, small gp120 peptides (5 to 20 amino acids in length, preferably 7 to 10 amino acids in length) or 15 peptidomimetics of gp120 that bind receptor, monoclonal antibodies that bind gp120, and small organic molecules that bind either gp120 or receptor.

The antibodies induced by the gp120 polypeptides can also be used to induce anti-idiotype antibodies 20 that bind CC chemokines. These anti-idiotype antibodies can be screened for binding to an anti-gp120 polypeptide antibody and inhibiting gp120 from binding CC-CKR receptor. Such anti-idiotype antibodies mimic gp120 by binding to CC-CKR receptor. Such antibodies, 25 preferably human antibodies, can be obtained in a number of ways, such as human antibodies from combinatorial libraries (e.g., Burton et al. *Adv. Immunol.* (1994) 57:191-280). It is now possible to produce transgenic animals (e.g., mice) that are 30 capable, upon immunization, of producing a full repertoire of human antibodies in the absence of endogenous immunoglobulin production. For example, homozygous deletion of the antibody heavy-chain joining 35 region (JH) gene in chimeric and germ-line mutant mice results in complete inhibition of endogenous antibody production. Transfer of the human germ-line

immunoglobulin gene array in such germ-line mutant mice results in the production of human antibodies upon antigen challenge as described in Jakobovitis et al., *Proc. Natl. Acad. Sci. USA* 90: 2551 (1993); Jakobovits et al., *Nature* 362:255-258 (1993); Bruggermann et al., *Year in Immuno.* 7: 33 (1993).

Alternatively, phage display technology as described by McCafferty et al., *Nature* 348:552-553 (1990) can be used to produce human antibodies and antibody fragments in vitro from immunoglobulin variable (V) domain gene repertoires from unimmunized donors. According to this technique, antibody V domain genes are cloned in-frame either into either a major or minor coat protein gene of a filamentous bacteriophage, such as M13 or fd, and displayed as functional antibody fragments on the surface of the phage particle. Because the filamentous particle contains a single-stranded DNA copy of the phage genome, selections based on the functional properties of the antibody also result in selection of the gene encoding the antibody exhibiting those properties. Phage display can be performed in a variety of formats as reviewed by, for example, Johnson, et al., *Current Opinion in Structural Biology* 3:564-571 (1993).

Several sources of V-gene segments can be used for phage display. Clackson et al., *Nature*, 352: 624-628 (1991) isolated a diverse array of anti-oxazolone antibodies from a small random combinatorial library of V genes derived from the spleens of immunized mice. A repertoire of V genes from unimmunized human donors (or embryonic cells) can be constructed. It has been demonstrated that antibodies to a diverse array of antigens (including self-antigens) can be isolated essentially following the techniques described by Marks et al., *J. Mol. Biol.*, 222: 581-597 (1991), or Griffith et al., *EMBO J.*, 12: 725-734 (1993).

In a natural immune response, antibody genes accumulate mutations at a high rate (somatic hypermutation). Some of the changes introduced confer higher affinity, and B cells displaying high-affinity surface immunoglobulin are preferentially replicated and differentiated during subsequent antigen challenge. This natural process can be mimicked by employing the technique known as "chain shuffling" (Marks et al., *Bio/Technol.* 10:779-783 [1992]). In this method, the affinity of "primary" human antibodies obtained by phage display can be improved by sequentially replacing the heavy and light chain V region genes with repertoires of naturally occurring variants (repertoires) of V domain genes obtained from unimmunized donors. This technique allows the production of antibodies and antibody fragments with affinities in the nM range. A strategy for making very large phage antibody repertoires has been described by Waterhouse et al., *Nucl. Acids Res.*, 21: 2265-2266 (1993).

Accordingly, antibodies that bind CC-CKR can be obtained by screening antibodies or fragments thereof expressed on the surface of bacteriophage in combinatorial libraries or in other systems as described above with a gp120 monoclonal antibody that inhibits gp120 binding to receptor.

In addition to screening antibodies with a gp-120 antibody, random or combinatorial peptide libraries can be screened with either a gp120 antibody or the gp120 compounds of the invention. Approaches are available for identifying peptide ligands from libraries that comprise large collections of peptides, ranging from 1 million to 1 billion difference sequences, which can be screened using monoclonal antibodies or target molecules. The power of this technology stems from the chemical diversity of the amino acids coupled with the

large number of sequences in a library. See for example, Scott et al., *Cur. Opin. Biotechnol.* 5(1):40-8 (1994); Kenan et al. *Trends Biochem. Sci.* (1994) 19(2):57-64. Accordingly, the monoclonal antibodies, 5 preferably human monoclonals or fragments thereof, generated as discussed herein, find use in treatment by inhibiting or treating HIV infection or disease progression, as well as in screening assays to identify additional pharmaceuticals.

10

#### Production of gp120

gp120 for a vaccine can be produced by any suitable means, as with prior art HIV gp120 subunit vaccines. Recombinantly-produced or chemically synthesized gp120 is preferable to gp120 isolated directly from HIV for safety reasons. Methods for recombinant production of gp120 are described below.

Oligonucleotides encoding gp120 from breakthrough isolates and capable of expressing gp120 can be prepared by conventional means. For example, the nucleotide sequence can be synthesized. Alternatively, another HIV nucleotide sequence encoding gp120 can be used as a backbone and altered at any differing residues as by site-directed mutagenesis. 25 Site-directed mutagenesis is described in Kunkel et al, *Proc. Natl. Acad. Sci. (USA)* 82:488-492 (1985) and Zoller et al, *Nuc. Acids Res.* 10:6487-6500 (1982) and is well known.

In a preferred embodiment, the nucleotide sequence 30 is present in an expression construct containing DNA encoding gp120 under the transcriptional and translational control of a promoter for expression of the encoded protein. The promoter can be a eukaryotic promoter for expression in a mammalian cell. In cases 35 where one wishes to expand the promoter or produce gp120 in a prokaryotic host, the promoter can be a

prokaryotic promoter. Usually a strong promoter is employed to provide high-level transcription and expression.

The expression construct can be part of a vector 5 capable of stable extrachromosomal maintenance in an appropriate cellular host or may be integrated into host genomes. Normally, markers are provided with the expression construct which allow for selection of a host containing the construct. The marker can be on 10 the same or a different DNA molecule, desirably, the same DNA molecule.

The expression construct can be joined to a replication system recognized by the intended host 15 cell. Various replication systems include viral replication systems such as those from retroviruses, simian virus, bovine papilloma virus, or the like. In addition, the construct may be joined to an amplifiable gene, e.g. the DHFR gene, so that multiple copies of the gp120 DNA can be made. Introduction of the 20 construct into the host will vary depending on the construct and can be achieved by any convenient means. A wide variety of prokaryotic and eukaryotic hosts can be employed for expression of the proteins.

Preferably, the gp120 is expressed in mammalian 25 cells that provide the same glycosylation and disulfide bonds as in native gp120. Expression of gp120 and fragments of gp120 in mammalian cells as fusion proteins incorporating N-terminal sequences of Herpes Simplex Virus Type 1 (HSV-1) glycoprotein D (gD-1) is 30 described in Lasky, L. A. et al., 1986 (Neutralization of the AIDS retrovirus by antibodies to a recombinant envelope glycoprotein) *Science* 233: 209-212 and Haffar, O.K. et al., 1991 (The cytoplasmic tail of HIV-1 gp160 contains regions that associate with cellular 35 membranes.) *Virol.* 180:439-441, respectively. A preferred method for expressing gp120 is described in

the examples. In the examples, a heterologous signal sequence was used for convenient expression of the protein. However, the protein can also be expressed using the native signal sequence.

5 An isolated, purified gp120 polypeptide having one of the amino acid sequences illustrated in Table 1 can be produced by conventional methods. For example, the proteins can be chemically synthesized. In a preferred embodiment, the proteins are expressed in mammalian 10 cells using an expression construct of this invention. The expressed proteins can be purified by conventional means. A preferred purification procedure is described in the examples.

15 gp120 Fragments

The present invention also provides gp120 fragments that are suitable for use in inducing antibodies for use in a vaccine formulation. A truncated gp120 sequence, as used herein, is a fragment 20 of gp120 that is free from a portion of the intact gp120 sequence beginning at either the amino or carboxy terminus of gp120. A truncated gp120 sequence of this invention is free from the C5 domain. The C5 domain of gp120 is a major immunogenic site of the molecule. 25 However, antibodies to the region do not neutralize virus. Therefore, elimination of this portion of gp120 from immunogens used to induce antibodies for serotyping is advantageous.

In another embodiment, the truncated gp120 30 sequence is additionally free from the carboxy terminal region through about amino acid residue 453 of the gp120 V5 domain. The portion of the V5 domain remaining in the sequence provides a convenient restriction site for preparation of expression 35 constructs. However, a truncated gp120 sequence that is free from the entire gp120 V5 domain is also

suitable for use in inducing antibodies.

In addition, portions of the amino terminus of gp120 can also be eliminated from the truncated gp120 sequence. In particular, the truncated gp120 sequence 5 can be free from the gp120 signal sequence. The truncated gp120 sequence can be free from the carboxy terminus through amino acid residue 111 of the gp120 C1 domain, eliminating most of the C1 domain but preserving a convenient restriction site. However, the 10 portion of the C1 domain through the V2 cysteine residue that forms a disulfide bond can additionally be removed, so that the truncated gp120 sequence is free from the carboxy terminus through amino acid residue 117 of the gp120 C1 domain. In a preferred embodiment, 15 the truncated gp120 sequence is free from the amino terminus of gp120 through residue 111 of the C1 domain and residue 453 through the carboxy terminus of gp120.

The truncated gp120 sequences can be produced by recombinant engineering, as described previously. 20 Conveniently, DNA encoding the truncated gp120 sequence is joined to a heterologous DNA sequence encoding a signal sequence.

It is understood that the application of the 25 teachings of the present invention to a specific problem or situation is within the capabilities of one having ordinary skill in the art in light of the teachings contained herein. Examples of the products of the present invention and representative processes for their isolation, use, and manufacture appear below, 30 but should not be construed to limit the invention.

All literature citations herein are expressly incorporated by reference.

35

#### EXAMPLES

##### Materials and Methods

Specimen collection from human volunteers. Blood was collected from MN-rgp120-immunized individuals who were infected with HIV-1 while participating in Phase I (NIH Protocol AVEG 016) and Phase II (NIH Protocol AVEG 201) HIV-1 vaccine trials sponsored by the National Institutes of Health (NIH). The demographics of the subjects in the study, and the study design have been described in McElrath; *Seminars in Cancer Biol.* 6:1-11 (1995); McElrath et al.; *Abstracts from Eighth Annual Meeting of the National Cooperative Vaccine Development Groups for AIDS*. Bethesda, MD 216 (1996). Specimens were obtained according to an informed consent protocol approved by the institutional review boards of the participating institutions. In the experimental section, the time of HIV-1 infection is specified with regard to data provided by the NIH AIDS Vaccine Evaluation Network where PCR (RNA) and/or serologic assays were used to detect HIV-1 infection.

Sample preparation for cloning HIV-1 envelope glycoproteins. Peripheral blood mononuclear cells (PBMCs) from HIV-1 infected vaccinees were prepared from heparinized venous blood by FICOLL-HYPAQUE gradient centrifugation. Cell number and viability were determined. After separation, PBMCs were washed twice in phosphate-buffered saline and suspended at a cell density of  $6 \times 10^6$  cells/ml in PCR lysis buffer (50 mM KCl, 10 mM Tris (pH 8.4), 2.5 mM MgCl<sub>2</sub>, 0.1 mg/ml gelatin (Sigma), 0.45% NONIDET P40 detergent, 0.45% TWEEN 20 detergent (both detergents are commercially available from United States Biochemical Corp.) and 0.06 mg/ml Proteinase K (Gibco BRL) to lyse the cells. The lysate was incubated at 50-60°C for 1 hour, followed by inactivation of the Proteinase K at 95°C for 10 minutes. Lysates were shipped frozen and stored at -70°C until use.

**Polymerase chain reaction (PCR) amplification.**

Samples were subjected to two rounds of PCR amplification using the nested primers described below. In the first round, 25  $\mu$ l aliquots of PBMC lysates (containing about 1  $\mu$ g genomic DNA) were mixed with an equal volume of a PCR reaction mix containing 400  $\mu$ M each dNTP, 200  $\mu$ g/ml BSA (Sigma Chemical Corporation, RIA grade) and about 100 pmoles of each primer in 50 mM KCl, 20 mM Tris (pH 8.4) and 3 mM MgCl<sub>2</sub>. After an initial 10 minute denaturation step at 95°C, 5 units of

10. Tag polymerase (AMPLITAQ, Perkin Elmer Cetus) were added during an 55°C soak step, and samples were overlayed with mineral oil.

The PCR profile was as follows: 2 cycles having 15. 1 minute at 55°C, 2.5 minutes at 72°C and 1 minute at 94°C, followed by 28 cycles with 30 seconds at 55°C, 2.5 minutes at 72°C and 45 seconds at 94°C, and an extension step at 72°C for 5 minutes.

Aliquots of 10  $\mu$ l from the first-round reactions 20. were re-amplified with appropriate nested primers in a final reaction volume of 100  $\mu$ l, using either the reagents and profile described above or the reagents and profile described in the PCR Optimizer Kit (Invitrogen.) PCR reaction products were purified 25. using QIAQUICK-spin columns (Qiagen Inc.) The primer pair used in the first round was either 120.os.F (5'-gggaattcgatccAGAGCAGAACAGTGGCAATGA with homologous sequence at position 6248-6270 of HIVPV22) (SEQ. ID. NO. 34) or JM11A 30. (5'-ctcgag-CTCCTGAAGACAGTCAGACTCATCAAG at position 6048-6074) (SEQ. ID. NO. 35) in the forward direction [Kusumi et al.; J. Virol. 66:875 (1992)] combined with 120.os.R (5'-ggcttagaagctttaGCCCATAGTGCTCCTGCTGCT-CC at position 7836-7859) (SEQ. ID. NO. 36) in the reverse 35. direction. The internal nested primers were 120.BX.F (5'-gggcggatcctcgaGGTACCTGTRTGGAAAGAAGCA at position

6389-6410; R: A or G) (SEQ. ID. NO. 37) and 120.is.R (5'-ggtctagaagtttaTGCTCCYAGAACCAAGGAACA at position 7819-7841; Y: T or C) (SEQ. ID. NO. 38). Heterologous primer sequences are shown in lower case letters.

5

**Subcloning of PCR products and the expression of recombinant envelope glycoproteins as fusion proteins.**

The HIV-1 envelope glycoprotein gp120 sequences were cloned and expressed as chimeric genes and fusion 10 proteins, where the signal sequence and 27 amino acids from the mature N terminus of herpes simplex virus type 1 (HSV-1) were fused to the N-terminal sequences of the gp120 genes, corresponding to amino acid 13 of the mature gp120 sequence. PCR products containing 15 gp120 sequences from the breakthrough specimens were cloned into pRK5 expression plasmid as chimeric genes using combinations of restrictions sites engineered into the heterologous PCR primer tails and the Xho I site engineered into the N-terminal sequence of 20 HSV-1 gD.

The resulting double-stranded DNA was sequenced with Sequenase and the dGTP Reagent Kit (United States Biochemical Corp.). Sequences from glycoprotein D were provided to enhance expression and to provide a flag 25 epitope to facilitate protein analysis, as described in Berman et al.; J. Virol. 7:4464-9 (1992); Nakamura et al.; AIDS and Human Retroviruses 8:1875-85 (1992); and Nakamura et al.; J. Virol. 67:6179-91 (1993).

Briefly, isolated DNA fragments generated by the 30 PCR reaction were ligated into a plasmid (pRK.gD-5, pRKgDstop) designed to fuse the gp120 fragments, in frame, to the 5' sequences of the glycoprotein D (gD) gene of Type 1 Herpes Simplex Virus (gD-1) and the 3' end to translational stop codons. The fragment of the 35 gD-1 gene encoded the signal sequence and 25 amino acids of the mature form of HSV-1 protein. To allow

for expression in mammalian cells, chimeric genes fragments were cloned into the pRK5 expression plasmid (Eaton et al., *Biochemistry* 291:8343-8347 (1986)) that contained a polylinker with cloning sites and 5 translational stop codons located between a cytomegalovirus promotor and a simian virus 40 virus polyadenylation site.

The resulting plasmids were transfected into the 293s embryonic human kidney cell line (Graham et al., 10 *J. Gen. Virol.* 36:59-77 (1977)) using a calcium phosphate technique (Graham et al., *Virology* 52:456-467 (1973)). Growth conditioned cell culture media was collected 48 hr after transfection, and the soluble proteins were detected by ELISA or by specific 15 radioimmunoprecipitation where metabolically labeled proteins from cell culture supernatants were resolved by sodium dodecyl sulfate polyacrylamide gel electrophoresis (PAGE) and visualized by autoradiography as described in Berman et al., 20 *J. Virol.* 63:3489-3498 (1989) and Laemmli, *Nature* 227:680-685 (1970).

**Serologic assays.** Sera were assayed for antibodies to rgp120, antibodies to synthetic gp120 V3 domain peptides corresponding to sequences from the gp120 V3 domain, and antibodies able to inhibit the binding of MN-rgp120 to cell surface CD4 using serologic assays described in Berman et al.; *J. Virol.* 7:4464-9 (1992); Nakamura et al.; *AIDS and Human Retroviruses* 8:1875-85 (1992); and Nakamura et al.; *J. Virol.* 67:6179-91 (1993). Endpoint titers of antibody binding to gp120 and V3 peptides were determined using three fold-serial dilutions of sera. The endpoint dilution titer was defined as the last 35 dilution that produced an optical density value that was two times higher than the mean of the optical

densities of 1:50 diluted, pooled, normal human sera. Antibody titers were calculated by a computer program that interpolated values between antibody dilutions. The inter-assay coefficient of variation of positive 5 control standard sera was 35%.

Binding of monoclonal antibodies to rgp120 from breakthrough viruses. An ELISA similar to that described by Moore et al.; AIDS 3:155-63 (1989) was 10 used to measure the binding of various monoclonal antibodies (MAbs) to rgp120s from breakthrough viruses. Briefly, Nunc-Immuno plates (Maxisorp, certified) were 15 coated (100  $\mu$ l at 5  $\mu$ g/ml in PBS at 4°C overnight) with an affinity-purified sheep polyclonal antiserum to a peptide at the C terminus of gp120 (D7324, International Enzymes, Fallbrook, CA). After washing once with PBS-0.05% TWEEN-20 detergent, the plates were 20 blocked with PBS-1.0% BSA for 30-60 minutes at room temperature. Cell culture supernatants from 293s cells, diluted to contain equivalent amounts of the 25 gD-rgp120 fusion protein, were added and incubated for 2 hours at room temperature followed by three washes with PBS-0.05% TWEEN-20 detergent. Various MAbs were diluted in PBS-1.0% BSA and 100  $\mu$ L of the diluted MAbs were added to each well and incubated for 1 hour at room temperature.

The plates were washed 3 times and incubated with 100  $\mu$ l of a horseradish peroxidase-conjugated second antibody (goat anti-mouse or anti-human IgG, Cappel) 30 for 1 hour at room temperature. After 3 washes the plates were developed and the OD<sub>492</sub> (optical density at 492 nm) read in a plate reader. Growth conditioned cell culture supernatants were normalized by dilution based on binding by MAb 5B6 which is specific for HSV-1 35 glycoprotein D fusion protein.

**Virus neutralization assays.** The ability of vaccinee sera to inhibit infection of MT4 cells by HIV-1<sub>MN</sub> was measured in a cytopathicity assay where cell viability was quantitated using a calorimetric indicator dye, as described in Robertson et al.; *J. Virol. Methods* 20:195-202 (1988). Briefly, a virus stock of HIV-1<sub>MN</sub> (obtained from Dr. Michael Norcross, U.S. Food and Drug Administration) was prepared as the clarified supernatant from chronically infected H9/HIV-1<sub>MN</sub> cell culture. H9 cells chronically infected with HIV-MN were pelleted and resuspended in one-tenth the original volume of medium. Cell-associated virus was released by the mechanical shearing effects of rapid vortexing of the cells as described in Wrin et al.; *J. Virol.* 69:39-48 (1995).

An amount of virus sufficient to ensure complete cell lysis killing in 7 days was incubated with three-fold serial dilutions of test antisera, and then used to challenge MT4 T-lymphoid cells in 10% FCS/RPMI-1640 cell culture media. The cultures were incubated for 7 days at 37°C in 5% CO<sub>2</sub>, and then cell viability was tested by the dye MTT, as described by Robertson et al.; *J. Virol. Methods* 20:195-202 (1988). Virus neutralization endpoints were quantitated by measurement of OD at 570-650 nm, and then the endpoint titers were calculated as the reciprocal of the antiserum dilution giving a signal that was two-fold above the control signal with unprotected (killed) cells. These titers were typically twice those calculated at 50% protection.

### Results

**Immunization history of infected subjects.** Since 1992, 499 adults have been immunized with MN-rgp120 in 35 Phase I trials in low or moderate risk individuals and in a Phase II clinical trial involving moderate to high

risk individuals. The studies described herein entail the genetic and immunologic characterization of the first seven of nine individuals who became infected with HIV-1 through high risk behavior during the course of these trials. A listing of the trials and summary of the status of the vaccinees is presented in Table 2A. A listing of the analysis of the vaccinees is presented in Table 2B.

10

TABLE 2A  
Description of Vaccinees Infected with HIV-1  
After Immunization with MN-rgp120

				*Antigen dose/
<u>Study No.</u>	<u>Case No.</u>	<u>*Risk Group</u>	<u>Adjuvant</u>	
15	016	C6	M/H	300/QS21
	016	C8	M/H	600/QS21
	016	C15	M/H	300/QS21
	201	C7	M/H	600/Alum
	201	C11	M/H	600/Alum
20	201	C10	M/IDU	600/Alum
	201	C17	M/IDU	600/Alum

\* - M/H indicates male homosexual; M/IDU indicate male intravenous drug user.

25                   † - numbers indicate dose in micrograms of MN-rgp120 injected per immunization; QS21 indicates antigen was formulated in QS21 adjuvant; Alum indicates MN-rgp120 formulated in aluminum hydroxide.

TABLE 2B  
Description of Vaccines Infected with HIV-1  
After Immunization with MN-rgp120

5	Case No.	Injection Schedule	Injections	Time of HIV-1+	■ Interval: to HIV-1+
		<u>(months)</u>	<u>HIV-1+</u>	<u>(months)</u>	<u>(months)</u>
	C6	0,1,10,5	2	4.00	2.00
	C8	0,1	2	4.00	3.00
	C15	0,1,2	3	6.25	4.00
10	C7	0,1,6,12	3	9.25	3.00
	C11	0,1,6,12	4	19.50	6.75
	C10	0,1,6,19	3	19.50	13.50
	C17	0,1,6,18	4	24.75	6.25

■ - indicates interval between last immunization  
and detection of HIV-1 infection.

Three of the infections occurred in a Phase I trial (NIH Protocol AVEG 201) that compared the safety and immunogenicity of MN-rgp120 formulated in two different adjuvants (alum and QS21), and four of the infections occurred in a Phase II trial aimed at establishing the safety and immunogenicity of MN-rgp120 in various high risk groups (e.g., intravenous drug users, homosexual and bisexual males, and partners of HIV-1 infected individuals).

Of the seven infections studied (Table 3), two (C6 and C8) occurred after two injections, three (C7, C10 and C15) occurred after three injections, and two (C11 and C17) occurred after receiving the four scheduled injections. The interval between receiving the last immunization and becoming infected was 2 to 13.5 months.

**TABLE 3**  
**Pak Pst Boost MN-rgp120 Antibody Titers**  
**in Vaccinees that Became Infected with HIV-1**

5	<b>Injections</b>	<b>C6</b>	<b>C8</b>	<b>C15</b>	<b>C7</b>	<b>C11</b>	<b>C10</b>	<b>C17</b>
	1	<50	2185	79	<50	1890	na	na
	2	21539	10125	na	413	32696	7771	7056
	3	#	#	4460	9707	34728	11627	1841
	4	#	#	#	#	#	#	1134
								0

10      # - indicates specimen not analyzed because of HIV-1 infection.

na - indicates the sample was not available for testing.

15      boldface - indicates unusually low antibody titers.

20      Antibody response to gp120 in vaccinated individuals. The magnitude and specificity of the antibody response to MN-rgp120 was measured by ELISA in all infected individuals throughout the course of the immunization regime (Figure 1). Five of the seven

subjects exhibited normal antibody response kinetics that included a small but reproducible primary response (1:100-1:2,000) and a strong secondary (booster) response (titers ranging from 1:7,000-1:32,000), and 5 antibody responses following third and fourth injections that were similar or marginally higher than those achieved after the second immunization (Figure 1, Table 3).

The antibody response observed in C7 (Figure 1C) 10 was unusual in that no antibodies were detectable after the primary injection and a titer of only 1:350 was detected after the second injection. It thus appeared that C7 did not respond to the primary immunization, and that the antibody response obtained after the 15 second injection represented a primary immune response. Consistent with this hypothesis, the third injection elicited a titer of only 1:9,707, typical of those normally seen after two immunizations.

An atypical antibody response was also seen in 20 subject C15 (Figure 1G) who was immunized according to an accelerated immunization schedule of 0, 1, and 2 months. As expected, the antibody titer seen in this subject (1:4,460) was at the low end of what is typically achieved after two immunizations and was far 25 below normal values for three immunizations. The lack of an effective booster response after the third immunization of C15 was not surprising in view of previous studies where an accelerated 0, 1, and 2 month 30 immunization schedule in baboons [Anderson et al.; *J. Infect. Dis.* 160:960-9 ((1989)) similarly prolonged the secondary response and failed to elicit an effective tertiary booster response.

Retrospective analysis of serum and plasma from 35 subjects C6 (Figure 1A) and C8 (Figure 1B) indicated that they became infected with HIV-1 at some point between the second and third immunizations. Serologic

evidence of HIV-1 infection was evident in the gp120 antibody assays where the titers failed to decline two weeks after the second injection and instead formed an uncharacteristic high titer plateau (Figures 1A and 1B). A similar plateau in MN-rgp120 titer after the third injection, suggested that subject C7 became infected around week 36, approximately 16 weeks after receiving the third injection (Figure 1C). Subjects C10 (Figure 1E), C11 (Figure 1D), C15 (Figure 1G), and 10 C17 (Figure 1F) developed unexpected increases in gp120 titers, typical of HIV-1 infection, after either the third or fourth immunizations. The data obtained demonstrate that immunologic priming for MN-rgp120 antibody responses is insufficient to provide universal 15 protection from HIV-1 infection.

**Antibody titers to the V3 domain.** To further characterize the antibody response to gp120, antibody titers were measured to a synthetic V3 domain peptide of MN-rgp120 containing the principal neutralizing determinant (PND). Five of the seven subjects developed good V3 titers (1:400 to 1:4000) after the second immunization, however two subjects (C7 and C15) required three immunizations before developing 25 significant titers (Figures 1C and 1G). As had been observed previously (11), the peak V3 titers in some individuals (e.g. C11, C10, C17) appeared to decline with each successive immunization (Figures 1D, 1E, and 1F). After HIV-1 infection, two patterns of V3 30 reactivity were observed. Three subjects (C6, C7, and C10) showed large increases in titer to V3 domain peptides (Figures 1A, 1C, and 1E) whereas C8 (Figure 1B) showed a large decrease in V3 titer. At the time of analysis, the data were insufficient to

draw any conclusions regarding the changes in V3 titers in response to HIV-1 infection in subjects C11, C15 and C17.

The results obtained indicate that the ability to 5 form antibodies reactive with the V3 domain at various time-points prior to HIV-1 infection is not a valid correlate of protective immunity against all strains of HIV-1.

10 **CD4 Inhibition titers.** Antibodies that block the binding of gp120 to CD4 represent a heterogeneous class of virus neutralizing antibodies. Some are known to bind to the C4 domain of gp120. [Nakamura et al.; *J. Virol.* 67:6179-91 (1993); Anderson et al.; *J. Infect. Dis.* 160:960-9 ((1989))], and some are known to 15 recognize conformation dependent discontinuous epitopes [Berman et al.; *J. Virol.* 7:4464-9 (1992); Nakamura et al.; *J. Virol.* 67:6179-91 (1993); McKeating et al.; *AIDS Research and Human Retroviruses* 8:451-9 (1992); Ho et al.; *J. Virol.* 65:489-93 (1991); 20 Barbas et al.; *Proc. Natl. Acad. Sci. USA* 91:3809-13 (1994)].

One way to detect antibodies to both types of epitopes is to measure the ability of vaccinee sera to 25 prevent the binding of [<sup>125</sup>I]-labeled gp120 to cell surface CD4 ([Nakamura et al.; *AIDS and Human Retroviruses* 8:1875-85 (1992); Nakamura et al.; *J. Virol.* 67:6179-91 (1993)]. CD4 blocking titers were detected in all seven of the vaccinees prior to 30 infection (Figure 2) with peak titers that ranged from 1:10-1:300. At the last time point prior to infection, the CD4 titers in five of the seven vaccinees was low (1:30 or less). One vaccinee (C17), however, possessed a CD4 blocking titer of about 1:300 prior to infection 35 (Figure 2F). Thus, the lack of antibodies that block the binding of MN-rgp120 to CD4 cannot account for all

of the infections. Large increases in CD4 blocking titers (1:100-1:1,000) were seen in five of the seven subjects after HIV-1 infection. These included vaccinees C6, C7, C8, C10, and C11. These results 5 demonstrate that the CD4 blocking titers elicited by MN-rgp120 were lower than those elicited by natural infection.

**Virus neutralizing activity.** The virus 10 neutralizing activity of antisera from MN-rgp120-immunized subjects was measured using a colorimetric assay that measured the viability of MT-4 cells after incubation with antibody treated virus (HIV-1<sub>MN</sub>). Since the actual date of infection was not 15 known for any of the breakthrough infections, and serum samples were collected infrequently, the magnitude of the neutralizing antibody response at the time of infection is not known for any of the vaccinees.

Of the seven infections examined, the serum sample 20 closest to the time of infection was that obtained from C7, where a neutralizing titer of 1:15 to HIV-1<sub>MN</sub> was present three weeks prior to detection of HIV-1 infection (Table 4). In all other cases, however, the 25 interval between the last injection and the time of infection was 10 to 25 weeks.

**TABLE 4**  
**Neutralization Activity of Sera from Vaccines**  
**Infected with HIV-1**

<u>Week</u>	<u>C6</u>	<u>C8</u>	<u>C15</u>	<u>C7</u>	<u>C11</u>	<u>C10</u>	<u>C17</u>
5 0	<10*	<10*	<10*	<10*	<10*	<10*	<10*
2	<10	<10	<10	-	-	-	-
4	<10*	<10*	nd*	<10*	<10*	<10*	<10*
6	10	80	-	<10	30	150	150
8	-	-	nd*	-	-	-	-
10	10	-	35	-	-	-	-
15	-	-	-	<10	-	-	-
16	150#	250#	-	-	30	10	<10
24			150#	<10*	20*	<10*	<10*
26				70	500	200	400
15 30				-	-	40	100
33				15	-	-	-
35				-	100	-	-
36				30#	-	10	40
52					30*	<10	<10
20 54						250	-
57						100	-
63						90	-
64						-	<10
77						40#	-
25 78							500# 10*
80							100
84							60

90

150

104

150#

5

\* - indicates immunization.

# - indicates HIV-1 positive.

nd - indicates not done.

-- indicates sample not available.

When sera from the two early infections were examined (Table 4), one individual (C6) had a peak neutralizing titer of 1:10 ten weeks prior to detection of HIV-1 infection, whereas the other individual (C8) had a neutralizing titer of 1:80 ten weeks prior to detection of HIV-1 infection. Subject C15, who was immunized according to an accelerated immunization schedule, developed a neutralizing titer of 1:35 after the third injection, 14 weeks prior to HIV-1 infection. Subject C10, who had a peak neutralizing titer of 1:200 following the third immunization (week 24), had no detectable titer at week 52, six months prior to the first indication of HIV-1 infection (week 78).

Subject C11 possessed a neutralizing titer of 1:90 at fourteen weeks prior to detection of HIV-1 and a peak titer of 1:500 following the third immunization.

Similarly vaccinee C17 had a neutralizing titer of 1:150 fourteen weeks prior to infection and a peak titer of 1:400 at two weeks after the third immunization.

Based on the rate of decay of the gp120 response of approximately two months [Belshe et al.; *JAMA* 272(6):475-80 (1994)], as well as the observation that neutralizing titers of 1:150 decayed to 1:10 in 10 weeks in vaccinees C10 and C17, it appears that neutralizing titers in C8, C15, C11, and C17 could have declined to 1:10 or less in the intervals between the last pre-infection serum sample and the time of HIV-1

detection.

The results of these studies demonstrated that all vaccinees developed some level of virus-neutralizing antibodies at some time prior to HIV-1 infection, and 5 that the magnitude of the neutralizing response was probably low at the time of infection. In general, the magnitude of the virus-neutralizing response observed in the individuals that became infected with HIV-1 was comparable to that seen in non-infected vaccinees as 10 described in Belshe et al.; *JAMA* 272(6):475-80 (1994).

Sequences of Viruses. To evaluate the similarity of the breakthrough viruses with the vaccine antigen, nucleotide sequences for gp120 from all seven 15 breakthrough viruses were determined. Envelope glycoprotein genes were amplified from proviral DNA using the polymerase chain reaction. Sequences were obtained by direct amplification of DNA from lysates of gradient-purified lymphocytes obtained directly from 20 patient blood without any intermediate tissue culture or amplification step.

A listing of the complete gp120 sequences (two clones per specimen) is provided in Figure 3. All 25 seven envelope glycoproteins possessed sequences typical of subtype (clade) B viruses. The overall homology with MN-rgp120 ranged from 69-80% (Table 5).

TABLE 5

Comparison of MN-rgp120 Sequence with Sequences  
from Infected Vaccinees\*

	MN	C6.1	C8.3	C7.2	C11.5	C10.5	C17.1	C15.2	
5	MN	100	79	78	70	75	69	80	72
	C6.1		100	78	70	81	75	90	79
	C8.3			100	68	80	76	84	83
	C7.2				100	80	73	76	73
	C11.5					100	75	70	80
10	C10.5						100	70	72
	C17.1							100	87
	C15.2								100

\* - Data indicate percent identity.

15 Interestingly, a high percentage (four of seven) of the breakthrough viruses differed from MN-rgp120 by 25-30% [Myers et al.; *Retroviruses and AIDS Database, Los Alamos National Laboratory* (1992 and 1995)]. Historically this degree of sequence variation is

20 typical of inter-subtype (intra-clade) variation rather than intra-subtype variation which is expected to be in the 10-20% range [Myers et al.; *Retroviruses and AIDS Database, Los Alamos National Laboratory* (1992 and 1995)]. Of the viruses with the greatest homology to

25 MN-rgp120, two (C6 and C8) occurred as early infections, prior to complete immunization, and one (C17) occurred as a late infection.

30 **Polymorphism in the V3 Domain.** Of particular interest were polymorphisms in regions known to contain epitopes recognized by virus neutralizing antibodies. The best characterized neutralizing epitope, the principal neutralizing determinant (PND), occurs at the

tip of the V3 loop. In subtype B viruses, approximately 60% possess the MN serotype-defining signature sequence, IGPGRAF (SEQ. ID. NO. 39), based on identity with the prototypic MN strain of HIV-1 [Berman et al.; *J. Virol.* 7:4464-9 (1992); Myers et al.; *Retroviruses and AIDS Database, Los Alamos National Laboratory* (1992 and 1995); La Rosa et al.; *Science* 249:932-5 (1990)].

Three of the viruses (C6, C8, and C17) possessed the MN serotype signature sequence (Figure 3). In contrast, four viruses possessed sequences with radical amino acid substitutions in the PND [IGPGRAW (C7), LGPGSTF (C11), IGPGRVL (C10), and IGPGSAT (C15)] (SEQ. ID. NOS. 40-43, respectively), and therefore were classified as "non-MN like" viruses. Of note, each of the four "non-MN-like" sequences were rare (Table 6) and were not typical of the most common "non-MN" variants of subtype B viruses [Myers et al.; *Retroviruses and AIDS Database, Los Alamos National Laboratory* (1992 and 1995)].

TABLE 6  
**Frequency of Polymorphisms at the Principal Neutralizing Determinant in HIV-1 Infected Individuals Immunized with MN-rgp120\***

5      V3 Sequence      Observed      Dataset Frequency

Sequence	n	Frequency (n=52)	GNE (n=52)	LANL (n=519)	LANL.1 (n=160)	LaRosa (n=245)
GPGRAF	3	0.42		0.67	0.57	0.66
GPGRAW	1	0.14		0.03	0.013	0.06
10      GPGRVL	1	0.14		<0.02	0.004	<0.006
GPGSTF**	1	0.14		<0.02	<0.002	<0.006
GPGSAF	1	0.14		0.02	0.011	<0.006

15      \* - Data set GNE refers to a collection of 52 independent isolates collected in 1992; dataset LANL refers to a collection of 519 sequences reported by Myers et al., *Retroviruses and AIDS Database, Los Alamos National Laboratory 1992 and 1995*; LANL.1 refers to a collection of 160 epidemiologically unlinked individuals provided by B. Korber (personal communication); dataset La Rosa refers to sequence data reported by La Rosa et al., *Science 249:932-5 (1990)*.

20      \*\* - Sequences were not present in the data sets examined.

30      The prevalence of viruses with PND sequences matching the breakthrough viruses ranged from a high of 1.3% (C7) to a low of 0.2% (C11) in a listing of 519 subtype B sequences compiled by the Los Alamos National Laboratory [Myers et al.; *Retroviruses and AIDS Database, Los Alamos National Laboratory (1992 and 1995)*]. Similarly low frequencies were observed in

three other independently derived data sets (Table 6). The occurrence of these sequences did not differ significantly between data sets collected prior to 1985 [La Rosa et al.; *Science* 249:932-5 (1990)] and data collected 1992, or from a set of 160 epidemiologically unlinked individuals (B. Korber, personal communication). All four sets of data agreed that the prevalence of viruses with MN-like PND sequences was in the range of 60%. Based on this data, four of the 10 seven breakthrough infections were determined to be caused by viruses that fell outside of the spectrum of viruses that the vaccine was expected to prevent.

**Other features of breakthrough virus V3 domains.**

15 Like MN-rgp120, the V3 domains of all of the breakthrough viruses were 36 amino acids in length. However, all seven viruses differed from MN-rgp120 with respect to the number of glycosylation sites and with respect to the syncytium-inducing (SI) signature 20 sequence.

The sequence of MN-rgp120 is somewhat unusual [Myers et al.; *Retroviruses and AIDS Database, Los Alamos National Laboratory* (1992 and 1995)] in that it lacks an N-linked glycosylation site at position 306 in 25 the V3 domain. The lack of this glycosylation site does not appear to be antigenically significant since antisera to MN-rgp120 are known to neutralize a variety of viruses (e.g. SF-2, DU6587-5, DU4489-5, CC) that possess a glycosylation site at this position 30 [Berman et al.; *J. Virol.* 7:4464-9 (1992)].

In addition, the V3 domain of MN-rgp120 possessed sequence polymorphisms (R at position 311, K at position 324, K at position 328) typical of syncytium inducing viruses [Fouchier et al.; *J. Virol.* 66:3183-87 35 (1992)], whereas all seven breakthrough viruses possessed sequences associated with non-syncytium-

inducing viruses. Syncytium-inducing viruses have been associated with rapid disease progression [Tersmette et al.; *J. Virol.* 62:2026-32 (1988)] and T cell tropism [O'Brien et al.; *Nature (London)* 348:69-73 (1990); 5 Shioda et al.; *Nature (London)* 349:167-9 (1991)]. To date viruses with these properties have not been recovered from any of the MN-rgp120 immunized volunteers.

10 **Polymorphism in the V1, V2 and C4 domains.**

Previous investigations have identified additional neutralizing epitopes in the V1, V2 and C4 domains of gp120 [Nakamura et al.; *J. Virol.* 67:6179-91 (1993); 15 McKeating et al.; *AIDS Research and Human Retroviruses* 8:451-9 (1992); Ho et al.; *J. Virol.* 65:489-93 (1991); Barbas et al.; *Proc. Natl. Acad. Sci. USA* 91:3809-13 (1994); McKeating et al.; *J. Virol.* 67:4932-44 (1993); Moore et al.; *J. Virol.* 67:6136-6151 (1993); Davis et al.; *J. Gen. Virol.* 74:2609-17 (1993)].

20 The best characterized of these neutralizing epitopes is in the C4 domain which has attracted special attention because antibodies binding to this area are known to block the binding of gp120 to CD4 [Moore et al.; *AIDS* 3:155-63 (1989); McKeating et al.; 25 *AIDS Research and Human Retroviruses* 8:451-9 (1992)]. Because the epitope is located in a conserved (C) domain, naturally-occurring polymorphism in this region is far more limited than in other neutralizing epitopes. Nakamura et al.; *J. Virol.* 67:6179-91 (1993) 30 reported that the binding of a number of neutralizing Mabs was dependent on K at position 429.

35 Comparison of the sequence of MN-rgp120 with other strains of HIV-1 showed that a common polymorphism, involving the substitution of E for K, occurs at this position. Indeed, substrains of the same virus isolate often show polymorphism at this position. The HXB2

substrain of HIV-1<sub>LA1</sub> contains K at position 429, whereas the BH10, IIIB, and LAV substrains of the HIV-1<sub>LA1</sub> contain E at this position [Nakamura et al.; J. Virol. 67:6179-91 (1993)]. Similarly, the 1984 5 isolate of HIV-1<sub>MN</sub> exhibited E at this position, while the 1990 isolate of HIV-1<sub>MN</sub>, used to produce MN-rgp120, possessed K at this position.

When the sequences of the infected vaccine 10 recipients were examined (Figure 3), the virus from subject C17, like MN-rgp120 contained K at position 429, whereas the six other viruses that differed from the vaccine immunogen possessed E at this position. These results demonstrated that six of the seven 15 breakthrough viruses differed from the vaccine immunogen at the CD4-blocking, neutralizing epitope in the C4 domain of gp120.

Studies with monoclonal antibodies have defined 20 neutralizing epitopes in the V1 and V2 domains of gp120 [McKeating et al.; J. Virol. 67:4932-44 (1993); Moore et al.; J. Virol. 67:6136-6151 (1993); Davis et al.; J. Gen. Virol. 74:2609-17 (1993)]. Like the 25 polymorphisms that occur in the C4 domain, the V2 domains exhibit several common polymorphisms that affect the binding of virus neutralizing antibodies. One such polymorphism occurs at position 171 which is 30 critically important for the binding of murine MAb 1025, whereas residue 187 is important for the binding of MAb several MAbs represented by 1088.

When the V2 domain sequences were examined 35 (Figure 3), all of the infected-vaccinee viruses differed from MN-rgp120 in that R replaced G at position 171 and I or V replaced E at position 187. Antibodies recognizing these adjacent sites in the V2 domain of MN-rgp120 would not be expected to neutralize viruses with radical amino acid substitutions at these position. Thus, all seven

breakthrough viruses differed from MN-rgp120 at a neutralizing epitope in the V2 domain of gp120.

Other neutralizing epitopes have been reported in the V1 domain of gp120 [O'Brien et al.; *Nature (London)* 5 348:69-73 (1990); McKeating et al.; *J. Virol.*

67:4932-44 (1993)]. Although the neutralizing epitopes in the V1 domain of MN-rgp120 have not been characterized, the polymorphism seen among the breakthrough viruses in this region was interesting.

10 Particularly striking (Figure 3) was that the length of this domain ranged from 20 amino acids (C17) to 45 amino acids (C6), and the number of N-linked glycosylation sites ranged from 2 to 6. In contrast, the V1 domain of MN-rgp120 is 31 amino acids in length 15 and encodes three N-linked glycosylation sites.

Although examination of sequence databases suggest that variation in the V2 region is comparable to the V1 region, the V2 region of the breakthrough viruses showed less variation than expected. Specifically, the 20 length of the V2 region ranged from 36 amino acids (C7) to 39 amino acids in length, with six of seven viruses containing three N-linked glycosylation sites in this domain. A high degree of polymorphism was found in the V4 region where sequences ranged from 26 (C10) to 33 25 (C15, C7) amino acids in length and contained either 4 or 5 N-linked glycosylation sites.

30 **Antigenicity of envelope glycoproteins from breakthrough viruses.** To determine the significance of sequence variation on glycoprotein antigenicity, recombinant gp120 was prepared from the viruses of all 35 seven infected vaccinees (Figure 4). In these studies a series of MAbs was assembled and their binding to MN-rgp120 was compared to that of rgp120 from the vaccinee isolates by ELISA (Table 7).

TABLE 7  
 Relative Reactivity\* of MAb Binding to rgp120 from  
 Infected Subjects Compared with Binding to MN-rgp120

		V3	Discontinuous	C8	V2
5	gp120	1034	50.1	1.5E	1025
	MN	1.0	1.00	1.00	1.00
	C6.1	0.37	0.37	0.17	0.00
	C6.5	0.33	0.33	0.75	0.00
	C8.3	0.11	0.37	0.38	0.00
10	C8.6	0.14	0.34	0.29	0.00
	C7.2	0.47	0.60	0.71	0.00
	C11.5	0.00	0.00	0.17	0.00
	C11.7	0.00	0.00	0.17	0.00
	C10.5	0.33	0.40	0.46	0.24
15	C10.7	0.42	0.48	0.50	0.29
	C17.1	0.33	0.52	0.33	0.00
	C17.3	0.37	0.56	0.33	0.00
	C15.2	0.00	0.47	0.92	0.00
	C15.3	0.00	0.37	0.63	0.00

20

\* - Relative reactivity values represent ratio of optical densities obtained with rgp120 from patient isolates divided by optical density obtained for MN-rgp120 at a MAb concentration of 2 micrograms per milliliter.

25

In control experiments, the binding of MAb 5B6 (which is specific for the HSV gD-1 flag epitope fused to the N terminus of all of the rgp120 protein) was used to standardize the amount of gp120 from each isolate (Figure 5A). These studies demonstrated that the assay was carried out under conditions where equivalent amount of rgp120s were captured onto wells of microtiter plates.

30

The antigenic structure of the V3 domain was examined using the 1034 MAb (isolated from mice immunized with MN-rgp120 as described in Nakamura et al.; J. Virol. 67:6179-91 (1993) and the 50.1 MAb (prepared from mice immunized with a synthetic V3 domain peptide as described in Rini et al.; Proc.

35

Natl. Acad. Sci. USA 90:6325-9 (1993). Both MAbs are known to exhibit potent virus neutralizing activity. When binding to the recombinant proteins was examined, the MAb binding to MN-rgp120 was at least 10-fold greater than to any of the breakthrough virus envelope proteins (Figure 5 B and C). Surprisingly, rgp120 from the three patient isolates (C8, C6, and C17) that possessed the MN serotype-defining sequence, IGPGR<sup>A</sup> (SEQ. ID. NO. 39), varied from one another in their MAb binding activity. Thus, the binding of MAb 1034 and MAb 50.1 to rgp120 from C17 was significantly greater than the binding to rgp120s from C6 and C8.

A distinction in the epitopes recognized by these MAbs was evident since C6-rgp120 and C8-rgp120 gave comparable binding with 50.1, whereas 1034 bound better to the C6-derived protein than the C8-derived protein. The poorest MAb reactivity was with rgp120s from C11 and C15. This result was consistent with sequence analysis demonstrating that these two viruses both possessed the radical substitution of S for R at position 18 in the V3 domain. Surprisingly, both of these MAbs exhibited better than expected binding to rgp120 from the C7 and C10 viruses. Like MN-rgp120, both proteins contained the penta-peptide, IGPGR<sup>A</sup> sequence (SEQ. ID. NO. 44) in the V3 loop, but differed from MN-rgp120 in that V and L replaced A and F at positions 319 and 320 in gp120 from C10, and W replaced F at position 320 in gp120 from C7. These results indicate that R at position 318 is essential for the binding of these two MAbs, and that the epitopes recognized by 1034 and 50.1 are not completely destroyed by the hydrophobic substitutions at positions 319 and 320.

As predicted from the sequence data, there was little if any binding to the breakthrough virus rgp120s using MAbs (1088 and 1025) directed to the V2 region of

MN-rgp120. Also consistent with sequence data was the observation that MAb 1024 directed to the C4 domain of MN-rgp120 gave some reactivity with C17-rgp120 which, like MN-rgp120 contained K at position 429, but gave no reactivity with the other isolates that contained E at residue 429.

Together, these studies demonstrated that the antigenic structure of all seven breakthrough viruses differed from the vaccine immunogen at three well characterized neutralizing epitopes.

A totally different pattern of reactivity was observed with the human hybridoma, MAb 15e, prepared from an HIV-1 infected individual as described in Ho et al.; *J. Virol.* 65:489-93 (1991). With this MAb, the greatest binding was achieved with MN-rgp120 and rgp120 from C7, and the poorest reactivity was seen with the two clones of rgp120 from the C11. Moderate, but comparable reactivity was seen with rgp120s from the C10 and C17.

These results demonstrate that the 15e epitope is polymorphic, and that the epitope is conserved on MN-rgp120 and rgp120 from C7, but has been lost on rgp120s from C11. Interestingly, the two different clones of gp120 derived from C6 gave strikingly different patterns of antibody binding. Thus, rgp120 from clone C6.5 exhibited strong reactivity with this antibody, whereas rgp120 from clones C6.1 exhibited significantly weaker activity with this MAb.

Comparison of sequence data (Figure 3) showed that the two C6 clones differed at 6 amino acid positions.

Based on comparative binding to the other viral proteins of known sequence, it appeared that the substitution of K for I at position 351 in the C3 domain of gp120 could account for the difference in binding activity. This result is also consistent with both clones of C11 similarly containing a positively-

charged K at this position, and also being poorly reactive with this MAb. Alternatively, a T for I substitution at position 439 in the C4 domain could account for the difference in 15e binding between C6.1 and C6.5. Although the inability of the two C11 clones to bind 15e cannot be explained by polymorphism at this position in the C4 domain, they could be affected by the adjacent T for M substitution at position 434.

10

#### Discussion

In these studies, the viruses and immune responses in seven of nine vaccinees who became infected with HIV-1 through high risk activity while participating in Phase I or Phase 2 trials of MN-rgp120, a candidate HIV-1 vaccine were analyzed. Such infections would be expected to occur for one of two reasons: 1) lack of sufficient immune response at the time of infection; or 2) infection with viruses that fall outside of the antigenic spectrum expected to be covered by the vaccine immunogen. The data indicate that both explanations may be involved with the infections observed (Table 8).

**TABLE 8**  
**Summary of Breakthrough Infections**  
Homologous to MN-rgp120

5	<u>Case No.</u>	<u>Adequate</u> <u>Immunization</u>	<u>Homology</u> (%)	<u>V3</u>	<u>C4</u>	<u>V2</u>	
						<u>PND</u>	<u>Epitope</u>
	C6	-	79	+	-	-	-
	C8	-	78	+	-	-	-
	C15	-	72	-	-	-	-
	C7	-	70	-	-	-	-
10	C11	+	75	-	-	-	-
	C10	+	69	-	-	-	-
	C17	+	80	+	+	-	-

15        Two of the infections occurred in individuals who failed to receive the minimum three doses of vaccine typically required for the induction of protective immunity with protein subunit vaccines (e.g. hepatitis B virus formulated in alum adjuvant as described in  
20        Francis et al.; *Ann. Int. Med.* 97:362-6 (1982)). Two additional breakthrough infections occurred in vaccinees who had weak or undetectable primary (C7) and booster (C15) responses. Of the three individuals who became infected with HIV-1 after receiving three or  
25        more productive immunizations (C10, C11, and C17), at least two, and possibly all three, appear to have become infected more than six months after receiving their last immunization. Because antibody titers to MN-rgp120 typically decay with a half-time of 2 to 2.5  
30        months [Belshe et al.; *JAMA* 272(6):475-80 (1994); Berman et al.; *AIDS* 8:591-601 (1994)], antibody titers would be expected to have decayed at least eight-fold and possibly as much as sixty four-fold at the time of infection. Thus, the lack of a sufficient immune

response at the time of infection represents a potential explanation for at least six of the seven breakthrough infections.

5 Data from vaccine efficacy studies in gp160 immunized chimpanzees [McElrath et al.; Longitudinal Vaccine-Induced Immunity and Risk Behavior of Study Participants in AVEG Phase II Protocol 201. In: Abstracts from Eighth Annual Meeting of the National Cooperative Vaccine Development Groups for AIDS.

10 Bethesda, MD 1996:216] challenged with HIV-1, and gp120-immunized rhesus macaques challenged with a chimeric SIV/HIV-1 virus (SHIV) suggest that the magnitude of the neutralizing antibody response at the time of infection is a critical correlate of protective immunity. If maintaining neutralizing antibody titers proves to be a valid correlate of protective immunity in humans, then formulations (e.g. novel adjuvants) or immunization regimes (frequent boosting) designed to maximize the antibody responses may be required to

15 achieve long lasting protection. Use of a booster every six months may be advantageous.

20

The other likely explanation for the late infections is the antigenic difference between the vaccine and the breakthrough virus envelope glycoproteins. This explanation is supported by the observation that four of the seven breakthrough viruses possessed envelope glycoproteins that differed from the MN-rgp120 by 25-30% at the amino acid level. Differences of this magnitude have historically 25 [Myers et al.; Retroviruses and AIDS Database, Los Alamos National Laboratory (1992 and 1995)] been 30 associated with inter-subtype variation and far exceeds the average 10-20% variation expected for viruses within the same subtype.

35 Although the biologic significance of sequence variation in many regions of the envelope glycoprotein

is unclear, polymorphism at neutralizing epitopes is an important factor that affects vaccine efficacy.

Previous studies [Salmon-Ceron et al.; *AIDS Res. and Human Retroviruses* 11:1479-86 (1995); Javaherian et al.; *Science* 250:1590-3 (1990)] have demonstrated that the breadth of neutralizing activity that could be elicited by HIV-1 envelope derived vaccines was critically dependent on the sequence of epitopes in the V3 domain (e.g.; the PND). Thus, candidate vaccines based on the LAI strain of HIV-1 (the prototypic "non-MN-like" subtype B virus), exhibited little or no cross neutralizing activity with subtype B viruses, whereas vaccines that contained the "MN-like" PND sequence (IGPGRAF) (SEQ. ID. NO. 44) exhibited broad cross neutralizing activity. That four of the seven breakthrough viruses possessed envelope glycoproteins with radical amino acid substitutions in the PND is consistent with the explanation that differences in antigenic structure explain some of these infections.

Over the last few years, it has become clear that polymorphism among "MN-like" viruses occurs at neutralizing epitopes outside of the PND. The best example occurs in the C4 domain where two antigenically distinct variants are distinguished by the presence of either K or E at position 429 [Moore et al.; *AIDS* 3:155-63 (1989)]. Because six of the seven breakthrough viruses differed from the vaccine strain in that they contained E rather than K at position 429, antibodies raised to the C4 domain of MN-rgp120 were unlikely to neutralize the viruses infecting in six of the seven vaccinees.

Other neutralizing epitopes are known to be present in the V1 and V2 domains of gp120. Although these regions are highly variable, due to insertions and deletions, neutralizing epitopes have been described by McKeating et al.; *J. Virol.* 67:4932-44

(1993); Moore et al.; *J. Virol.* 67:6136-6151 (1993); and Davis et al.; *J. Gen. Virol.* 74:2609-17 (1993). Several of these epitopes overlap an amino terminal sequence of the V2 domain containing the tri-peptide sequence RDK at positions corresponding to 142 to 144 of MN-rgp120 [McKeating et al.; *J. Virol.* 67:4932-44 (1993); Moore et al.; *J. Virol.* 67:6136-6151 (1993)]. Like the C4 epitope, variation in this sequence is known to occur between different substrains derived from the same parental isolate. Since all seven breakthrough viruses differed from MN-rgp120 in that they possessed the RDK sequence, rather than the GDK sequence present in the vaccine antigen, neutralizing antibodies to the V2 domain of MN-rgp120 would not have been expected neutralize any of the viruses recovered from the vaccinees immunized with MN-rgp120.

Although polymorphisms at neutralizing epitopes might account for the lack of protection in most of the infections, this does not appear to explain the infection of vaccinee C17, who was infected by a virus that matched MN-rgp120 in the V3 and C4 domains. If a difference in sequence was responsible for the lack of protection in this case, the critical difference might relate to the unusual sequence in the V1 domain of gp120 from this breakthrough virus. Several studies have shown that the V1 domain possesses epitopes recognized by virus neutralizing monoclonal antibodies [McKeating et al.; *J. Virol.* 67:4932-44 (1993); Davis et al.; *J. Gen. Virol.* 74:2609-17 (1993); Kayman et al.; *J. Virol.* 68:400-410 (1994)].

Although far less is known about the V1 epitopes relative to other neutralizing sites, the V1 epitopes appear to be conformation-dependent, and antisera from HIV-1 infected individuals recognize epitopes in the V1 and V2 domains [McKeating et al.; *J. Virol.* 67:4932-44 (1993); Kayman et al.; *J. Virol.* 68:400-410 (1994)].

The V1 sequence of the virus from C17 is noteworthy because it is smaller and contains fewer N-linked glycosylation sites than that of MN-rgp120 or any of the other breakthrough viruses. By the same token, the 5 envelope glycoproteins from C11 and C6 are noteworthy because they are significantly larger and contain more glycosylation sites than MN-rgp120 or the other breakthrough viruses.

While differences in amino acid sequence can 10 provide clues to differences in antigenic structure, the consequences of such polymorphism can only be proven through antibody binding studies. To correlate differences in sequence with differences in antigenic structure, gp120 from two clones each of all seven 15 breakthrough viruses was expressed and the antigenicity of the clones with a panel of monoclonal antibodies was examined. As predicted from the sequence data, none of the breakthrough virus envelope glycoproteins reacted with neutralizing MAbs to the V2 domain of MN-rgp120. 20 When MAbs to the C4 domain were examined, only the C17 envelope glycoprotein (that matched MN-rgp120 with respect to K429) showed significant, albeit lower, binding. Surprisingly, the three breakthrough envelope glycoproteins that contained the subtype B PND 25 consensus sequence, IGPGRAY (SEQ. ID. NO. !!), gave poor reactivity with all three PND directed MAbs, even though they possessed PND sequences closely related to the vaccine immunogen. Thus, all three of the vaccinee isolates appeared to possess changes outside of the 30 recognition site that interfered with MAb binding.

It has been known for many years that resistance to neutralization in vitro can sometimes be attributed to mutations in remote sequences that alter the conformation of neutralizing epitopes and interfere 35 with recognition by virus neutralizing antibodies [Nara et al.; *J. Virol.* 64:3779-91 (1990); Cordonnier

et al.; *Nature* 340:571-4 (1989)]. Together, these results indicate that the antigenic structure of the envelope glycoproteins recovered from the breakthrough viruses differed significantly from that of the vaccine antigen.

5 A novel result was the localization of residues in the C3 domain that appeared to affect the binding of the virus neutralizing human MAb, 15e. This MAb is known to recognize a discontinuous epitope, block CD4 10 binding, and neutralize a variety of laboratory and primary isolates of HIV-1 [Ho et al.; *J. Virol.* 65:489-93 (1991); Thali et al.; *J. Virol.* 66:5635-5641 (1992); Moore et al.; *AIDS Res. Hum. Retroviruses* 9:1179-1187 (1993)].

15 Comparative binding to envelope glycoproteins from the breakthrough viruses indicated that recognition by this antibody is critically dependent on residues in the C3 or C4 domains of gp120. The unique occurrence of a positively charged K at position 351 in the C3 20 domain provides a common explanation for the inability of the C11.5, C11.7 and C6.1 strains of HIV-1 to bind to 15e. Alternatively, it is possible that different amino acid substitutions in different locations account for the failure of 15e to bind to rgp120s from the C6 25 and C11 clones. The only obvious positions where substitutions of this type occur are in the C4 domain where T replaces M at 434 (C11) and T replaces I at 439.

30 The present studies demonstrate that the current formulation of MN-rgp120 is less than 100% effective against HIV-1 infection. Based on previous in vitro and in vivo studies with MN-rgp120, protection from natural HIV-1 infection in humans is expected to depend on a threshold concentration of virus-neutralizing 35 antibodies, and antigenic similarity between the vaccine immunogen and the challenge virus.

In this regard, only one of the seven breakthrough infections (C17) was unexpected. This individual received a full course of immunizations yet became infected with a virus similar to MN-rgp120 at least

5. two important neutralizing epitopes (V3 and C4 domains). This infection might be related to the magnitude of the antibody response at the time of infection, or antigenic differences between the breakthrough virus and the vaccine strain, or  
10. circumstances of infection (e.g., ulcerative lesions, infection by donor with acute infection or high viremia), not monitored in this protocol.

Alternatively this individual may represent a true vaccine failure, without clear explanation.

15. On balance, the analysis of breakthrough infections described herein did not uncover any data that would discourage the continued development of MN-rgp120 as a vaccine to prevent HIV-1 infection. The results support speculation that enhancing vaccine  
20. immunogenicity (as by additional booster immunizations) may be required to maintain long term protective immunity, and that the addition of rgp120 from other antigenically different strains of virus in addition to MN-rgp120 are useful to expand the breadth of  
25. protection.

The availability of viruses and viral glycoproteins derived from breakthrough infections may provide an important means to streamline the process of identifying new antigens for inclusion into a  
30. multivalent vaccine. Recombinant viral glycoproteins prepared from breakthrough viruses, by definition, possess antigenic structures that are significantly different from MN-rgp120, and are representative of viruses currently being transmitted. Thus, combining  
35. rgp120 from breakthrough viruses with MN-rgp120 is an effective way complement and significantly expand

**antigenic complexity and increase breadth of cross neutralizing activity.**

## SEQUENCE LISTING

(1) GENERAL INFORMATION:

5 (i) APPLICANT: Berman, Phillip W.

(ii) TITLE OF INVENTION: HIV ENVELOPE POLYPEPTIDES AND  
VACCINE

(iii) NUMBER OF SEQUENCES: 44

(iv) CORRESPONDENCE ADDRESS:

10 (A) ADDRESSEE: SKJERVEN, MORRILL, MACPHERSON, ET AL.

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(D) STATE: California

(E) COUNTRY: USA

(F) ZIP: 95110

15 (v) COMPUTER READABLE FORM:

(A) MEDIUM TYPE: 3.5 inch, 1.44 Mb floppy disk

(B) COMPUTER: IBM PC compatible

(C) OPERATING SYSTEM: PC-DOS/MS-DOS

(D) SOFTWARE: WinPatin (Genentech)

20 (vi) CURRENT APPLICATION DATA:

(A) APPLICATION NUMBER:

(B) FILING DATE:

(C) CLASSIFICATION:

(viii) ATTORNEY/AGENT INFORMATION:

25 (A) NAME: Terlizzi, Laura

(B) REGISTRATION NUMBER: 31,307

(C) REFERENCE/DOCKET NUMBER: M-3897 US

(ix) TELECOMMUNICATION INFORMATION:

30 (A) TELEPHONE: (408) 453-9200

(B) TELEFAX: (408) 453-7979

(2) INFORMATION FOR SEQ ID NO:1:

(i) SEQUENCE CHARACTERISTICS:

35 (A) LENGTH: 1503 base pairs

(B) TYPE: Nucleic Acid

(C) STRANDEDNESS: Single

(D) TOPOLOGY: Linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:1:

40 GGG GTA CCT GTG TGG AAG GAA GCA ACC ACC ACT CTA 36  
Gly Val Pro Val Trp Lys Glu Ala Thr Thr Thr Leu  
1 5 10

45 TTT TGT GCA TCA GAT GCT AAA GCA TAT GAC ACA GAG GTG 75  
Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu Val  
15 20 25

50 CAT AAT GTT TGG GCC ACA CAT GCT TGT GTA CCC ACA GAC 114  
His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp  
30 35

55 CCA AAC CCA CAA GAA ATG GTA TTG GAA AAT GTG ACA GAA 153  
Pro Asn Pro Gln Glu Met Val Leu Glu Asn Val Thr Glu  
40 45 50

60 GAT TTT AAC ATG TGG AAA AAT GAC ATG GTA GAA CAG ATG 192  
Asp Phe Asn Met Trp Lys Asn Asp Met Val Glu Gln Met  
55 60

65 CAT GAG GAT ATA ATC AGT TTA TGG GAT CAA AGC CTA AAA 231  
His Glu Asp Ile Ile Ser Leu Trp Asp Gln Ser Leu Lys  
65 70 75

CCA TGT CTA AAA TTA ACC CCA CTC TGT ATT ACT TTA AAT 270  
 Pro Cys Val Lys L u Thr Pro Leu Cys Ile Thr Leu Asn  
 80 85 90

5 TGC ACC AAT TGG AAG AAG AAT GAT ACT AAA ACT AAT AGT 309  
 Cys Thr Asn Trp Lys Lys Asn Asp Thr Lys Thr Asn Ser  
 95 100

10 AGT AGT ACT ACA ACT AAT AAT AGT AGT GCT ACA GCT AAT 348  
 Ser Ser Thr Thr Asn Asn Ser Ser Ala Thr Ala Asn  
 105 110 115

15 AGT AGT AGT ACT ACA ACT AAT AGT AGT TGG GGA GAG ATA 387  
 Ser Ser Thr Thr Asn Ser Ser Trp Gly Glu Ile  
 120 125

20 AAG GAG GGA GAA ATA AAG AAC TGC TCT TTC AAT ATC ACC 426  
 Lys Glu Gly Glu Ile Lys Asn Cys Ser Phe Asn Ile Thr  
 130 135 140

25 ACA AGC ATA AGA GAC AAG GTG AAG AAA GAA TAT GCA CTT 465  
 Thr Ser Ile Arg Asp Lys Val Lys Lys Glu Tyr Ala Leu  
 145 150 155

30 TTT TAT AGC CTT GAT GTA GTA CCA ATA GAA AAT GAT AAT 504  
 Phe Tyr Ser Leu Asp Val Val Pro Ile Glu Asn Asp Asn  
 160 165

35 ACT AGC TAT AGG TTG AGA AGT TGT AAC ACC TCA GTC ATT 543  
 Thr Ser Tyr Arg Leu Arg Ser Cys Asn Thr Ser Val Ile  
 170 175 180

40 ACA CAA GCC TGT CCA AAG GTA ACT TTT GAG CCA ATT CCC 582  
 Thr Gln Ala Cys Pro Lys Val Thr Phe Glu Pro Ile Pro  
 185 190

45 ATA CAT TAT TGT ACC CCG GCT CGT TTT CGC ATT CTG AAG 621  
 Ile His Tyr Cys Thr Pro Ala Gly Phe Ala Ile Leu Lys  
 195 200 205

50 TGT AGA GAT AAA AAG TTC AAT GGA ACA GGA CCA TGC AAA 660  
 Cys Arg Asp Lys Phe Asn Gly Thr Gly Pro Cys Lys  
 210 215 220

55 AAT GTT AGC ACA GTA CAA TGT GCA CAT GGA ATT AAG CCA 699  
 Asn Val Ser Thr Val Gln Cys Ala His Gly Ile Lys Pro  
 225 230

60 GTA GTG TCA ACT CAA CTG CTG TTA AAT GGC AGC CTA GCA 738  
 Val Val Ser Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala  
 235 240 245

65 GAA GAA GAG GTA ATA ATT AGA TCT GCC AAT TTC TCA AAC 777  
 Glu Glu Glu Val Ile Ile Arg Ser Ala Asn Phe Ser Asn  
 250 255

70 AAT GCT AAA ATC ATA ATA GTA CAG TTG AGG GAA CCT GCA 816  
 Asn Ala Lys Ile Ile Val Gln Leu Arg Glu Pro Val  
 260 265 270

75 GAA ATT AAT TGT ACA AGA CCC AGC AAC AAT ACA ATA AAA 855  
 Glu Ile Asn Cys Thr Arg Pro Ser Asn Asn Thr Ile Lys  
 275 280 285

GGT ATA CAC ATA CGA CCA GGG AGA GCA TTT TAT GCA ACA 894  
 Gly Ile His Ile Gly Pro Gly Arg Ala Phe Tyr Ala Thr  
 290 295

5 GGA GAC ATA CGA CGA GAT ATA AGA CAA GCA CAT TGT AAC 933  
 Gly Asp Ile Arg Gly Asp Ile Arg Gln Ala His Cys Asn  
 300 305 310

ATT AGT CGA GCA AAA TGG AAT AAC ACT TTA AAG AAG GTA 972  
 10 Ile Ser Gly Ala Lys Trp Asn Asn Thr Leu Lys Lys Val  
 315 320

GTT AAA AAA TTA AAA GAA CAA TTT CCA AAT AAA ACA ATA 1011  
 Val Lys Lys Leu Lys Glu Gln Phe Pro Asn Lys Thr Ile  
 15 325 330 335

GTC TTT AAC CAT TCC TCA GGA GGG GAC CCA GAA ATT GTA 1050  
 Val Phe Asn His Ser Ser Gly Gly Asp Pro Glu Ile Val  
 340 345 350

20 ATG CAC AGT TTT AAT TGT CAA GGG GAA TTT TTC TAC TGT 1089  
 Met His Ser Phe Asn Cys Gln Gly Glu Phe Phe Tyr Cys  
 355 360

25 AAT ACA ACA AAG CTG TTT AAT AGT ACT TGG AAT GAT ACT 1128  
 Asn Thr Thr Lys Leu Phe Asn Ser Thr Trp Asn Asp Thr  
 365 370 375

30 ACA GAG TCA AAT AAC AAT GAT AGT ACT ATT ACA CTC CCA 1167  
 Thr Glu Ser Asn Asn Asn Asp Ser Thr Ile Thr Leu Pro  
 380 385

TGC AGA ATA AAA CAA ATT ATA AAC ATG TGG CAG GAA ATA 1206  
 Cys Arg Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Ile  
 390 395 400

35 GGA AAA GCA ATG TAT GCC CCT CCC ACC AGA GGA GAA ATT 1245  
 Gly Lys Ala Met Tyr Ala Pro Pro Thr Arg Gly Glu Ile  
 405 410 415

40 AAA TGT TCA TCA AAT ATT ACA GGA CTA CTG TTA ATA AGA 1284  
 Lys Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Ile Arg  
 420 425

45 GAT GGT GGT ATT AAC ACT AGC GAT GCC ACC GAG ACC TTC 1323  
 Asp Gly Gly Ile Asn Thr Ser Asp Ala Thr Glu Thr Phe  
 430 435 440

50 AGA CCG GGA GGA GGA GAT ATG AGG GAC AAT TGG AGA ACT 1362  
 Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp Arg Ser  
 445 450

GAA TTA TAT AAA TAT AAA GTA GTG AAA ATT GAG CCA TTA 1401  
 Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile Glu Pro Leu  
 455 460 465

55 GGA GTA GCA CCC ACC AAAG GCA AAG AGA AGA GTG GTG CAG 1440  
 Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln  
 470 475 480

60 AGA GAA AAA AGA GCA GTA ACA CTA GGA GCT ATG TTC CTT 1479  
 Arg Glu Lys Arg Ala Val Thr Leu Gly Ala Met Phe Leu  
 485 490

GGG TTC TTA GGA GCA TAA AGC TTC 1503  
 Gly Phe Leu Gly Ala Xaa Ser Phe  
 495 500 501

## 5 (2) INFORMATION FOR SEQ ID NO:2:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 501 amino acids
- (B) TYPE: Amino Acid
- (C) TOPOLOGY: Linear

## 10 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:2:

Gly Val Pro Val Trp Lys Glu Ala Thr Thr Thr Leu Phe Cys Ala  
 1 5 10 15

15 Ser Asp Ala Lys Ala Tyr Asp Thr Glu Val His Asn Val Trp Ala  
 20 25 30

Thr His Ala Cys Val Pro Thr Asp Pro Asn Pro Gln Glu Met Val  
 35 40 45

20 Leu Glu Asn Val Thr Glu Asp Phe Asn Met Trp Lys Asn Asp Met  
 50 55 60

Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp Asp Gln Ser  
 65 70 75

Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Ile Thr Leu Asn  
 80 85 90

30 Cys Thr Asn Trp Lys Lys Asn Asp Thr Lys Thr Asn Ser Ser Ser  
 95 100 105

Thr Thr Thr Asn Asn Ser Ser Ala Thr Ala Asn Ser Ser Ser Thr  
 110 115 120

35 Thr Thr Asn Ser Ser Trp Gly Glu Ile Lys Glu Gly Glu Ile Lys  
 125 130 135

40 Asn Cys Ser Phe Asn Ile Thr Thr Ser Ile Arg Asp Lys Val Lys  
 140 145 150

Lys Glu Tyr Ala Leu Phe Tyr Ser Leu Asp Val Val Pro Ile Glu  
 155 160 165

45 Asn Asp Asn Thr Ser Tyr Arg Leu Arg Ser Cys Asn Thr Ser Val  
 170 175 180

Ile Thr Gln Ala Cys Pro Lys Val Thr Phe Glu Pro Ile Pro Ile  
 185 190 195

50 His Tyr Cys Thr Pro Ala Gly Phe Ala Ile Leu Lys Cys Arg Asp  
 200 205 210

55 Lys Lys Phe Asn Gly Thr Gly Pro Cys Lys Asn Val Ser Thr Val  
 215 220 225

Gln Cys Ala His Gly Ile Lys Pro Val Val Ser Thr Gln Leu Leu  
 230 235 240

60 Leu Asn Gly Ser Leu Ala Glu Glu Val Ile Ile Arg Ser Ala  
 245 250 255

Asn Phe Ser Asn Asn Ala Lys Ile Ile Val Gln Leu Arg Glu  
 260 265 270

Pro	Val	Glu	Ile	Asn	Cys	Thr	Arg	Pro	Ser	Asn	Asn	Thr	Ile	Lys		
275															285	
Gly	Ile	His	Ile	Gly	Pro	Gly	Arg	Ala	Phe	Tyr	Ala	Thr	Gly	Asp		
5															290 295 300	
Ile	Arg	Gly	Asp	Ile	Arg	Gln	Ala	His	Cys	Asn	Ile	Ser	Gly	Ala		
															305 310 315	
10	Lys	Trp	Asn	Asn	Thr	Leu	Lys	Lys	Val	Val	Lys	Lys	Leu	Lys	Glu	
															320 325 330	
Gln	Phe	Pro	Asn	Lys	Thr	Ile	Val	Phe	Asn	His	Ser	Ser	Gly	Gly		
15															335 340 345	
Asp	Pro	Glu	Ile	Val	Met	His	Ser	Phe	Asn	Cys	Gln	Gly	Glu	Phe		
															350 355 360	
Phe	Tyr	Cys	Asn	Thr	Thr	Lys	Leu	Phe	Asn	Ser	Thr	Trp	Asn	Asp		
20															365 370 375	
Thr	Thr	Glu	Ser	Asn	Asn	Asn	Asp	Ser	Thr	Ile	Thr	Leu	Pro	Cys		
															380 385 390	
25	Arg	Ile	Lys	Gln	Ile	Ile	Asn	Met	Trp	Gln	Glu	Ile	Gly	Lys	Ala	
															395 400 405	
Met	Tyr	Ala	Pro	Pro	Thr	Arg	Gly	Glu	Ile	Lys	Cys	Ser	Ser	Asn		
															410 415 420	
30	Ile	Thr	Gly	Leu	Leu	Leu	Ile	Arg	Asp	Gly	Gly	Ile	Asn	Thr	Ser	
															425 430 435	
Asp	Ala	Thr	Glu	Thr	Phe	Arg	Pro	Gly	Gly	Gly	Asp	Met	Arg	Asp		
35															440 445 450	
Asn	Trp	Arg	Ser	Glu	Leu	Tyr	Lys	Tyr	Lys	Val	Val	Lys	Ile	Glu		
															455 460 465	
40	Pro	Leu	Gly	Val	Ala	Pro	Thr	Lys	Ala	Lys	Arg	Arg	Val	Val	Gln	
															470 475 480	
Arg	Glu	Lys	Arg	Ala	Val	Thr	Leu	Gly	Ala	Met	Phe	Leu	Gly	Phe		
															485 490 495	
45	Leu	Gly	Ala	Xaa	Ser	Phe										
															500 501	

## (2) INFORMATION FOR SEQ ID NO:3:

## (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1503 base pairs

(B) TYPE: Nucleic Acid

(C) STRANDEDNESS: Single

(D) TOPOLOGY: Linear

## 55 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:3:

GGG	GTA	CCT	GTA	TGG	AAA	GAA	GCA	ACC	ACC	ACT	CTA	36
Gly	Val	Pro	Val	Trp	Lys	Glu	Ala	Thr	Thr	Thr	Leu	

1					5						10	
---	--	--	--	--	---	--	--	--	--	--	----	--

60	TTT	TGT	GCA	TCA	GAT	GCT	AAA	GCA	TAT	GAC	ACA	GAG	GTG	75
	Phe	Cys	Ala	Ser	Asp	Ala	Lys	Ala	Tyr	Asp	Thr	Glu	Val	

15									20				25	
----	--	--	--	--	--	--	--	--	----	--	--	--	----	--

CAT AAT GTT TGG GCC ACA CAT GCT TGT GTA CCC ACA GAC 114  
 His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp  
 30 35

5 CCA AAC CCA CAA GAA ATG GTA TTG GAA AAT GTG ACA GAA 153  
 Pro Asn Pro Gln Glu Met Val Leu Glu Asn Val Thr Glu  
 40 45 50

10 GAT TTT AAC ATG TGG AAA AAT GAC ATG GTA GAA CAG ATG 192  
 Asp Phe Asn Met Trp Lys Asn Asp Met Val Glu Gln Met  
 55 60

CAT GAG ANT ATA ATC AGT TTA TGG GAT CAA AGC CTA AAA 231  
 His Glu Xaa Ile Ile Ser Leu Trp Asp Gln Ser Leu Lys  
 15 65 70 75

CCA TGT GTA AAA TTA ACC CCA CTC TGT ATT ACT TTA AAT 270  
 Pro Cys Val Lys Leu Thr Pro Leu Cys Ile Thr Leu Asn  
 80 85 90

20 TGC ACC AAT TGG AAG GAG AAT GAT ACT AAA ACT AAT AGT 309  
 Cys Thr Asn Trp Lys Glu Asn Asp Thr Lys Thr Asn Ser  
 95 100

25 AGT AGT ACT ACA ACT AAT AGT AGT GCT ACA GCT AAT 348  
 Ser Ser Thr Thr Asn Asn Ser Ser Ala Thr Ala Asn  
 105 110 115

30 AGT AGT AGT ACT ACA ACT AAT AGT AGT TGG CGA GAC ATA 387  
 Ser Ser Ser Thr Thr Asn Ser Ser Trp Gly Glu Ile  
 120 125

35 AAG GAG GGA GAA ATA AAG AAC TGC TCT TTC AAT ATC ACC 426  
 Lys Glu Gly Glu Ile Lys Asn Cys Ser Phe Asn Ile Thr  
 130 135 140

ACA GGC ATA AGA GAC AAG GTG AAG AAA GAA TAT GCA CTT 465  
 Thr Gly Ile Arg Asp Lys Val Lys Lys Glu Tyr Ala Leu  
 145 150 155

40 TTT TAT AGC CTT GAT GTA GTA CCA ATA GAA AAT GAT AAT 504  
 Phe Tyr Ser Leu Asp Val Val Pro Ile Glu Asn Asp Asn  
 160 165

45 ACT AGC TAT AGG TTG AGA AGT TGT AAC ACC TCA GTC ATT 543  
 Thr Ser Tyr Arg Leu Arg Ser Cys Asn Thr Ser Val Ile  
 170 175 180

50 ACA CAA GCC TGT CCA AAG GTA ACT TTT GAG CCA ATT CCC 582  
 Thr Gln Ala Cys Pro Lys Val Thr Phe Glu Pro Ile Pro  
 185 190

55 ATA CAT TAT TGT ACC CCG GCT GGT TTT GCG ATT CTG AAG 621  
 Ile His Tyr Cys Thr Pro Ala Gly Phe Ala Ile Leu Lys  
 195 200 205

TGT AAA GAT AAA AAG TTC AAT GGA ACA GGA CCA TCC AAA 660  
 Cys Lys Asp Lys Lys Phe Asn Gly Thr Gly Pro Cys Lys  
 210 215 220

60 AAT GTT AGC ACA GCA CAA TGT ACA CAT GGA ATT AAG CCA 699  
 Asn Val Ser Thr Val Gln Cys Thr His Gly Ile Lys Pro  
 225 230

235      GTA GTG TCA ACT CAA CTG CTG TTA AAT GGC AGC CTA GCA 738  
 Val Val Ser Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala  
 240      245  
 5      GAA GAA GAG GTA ATA ATT AGA TCT GCC AAT TTC TCA AAC 777  
 Glu Glu Glu Val Ile Ile Arg Ser Ala Asn Phe Ser Asn  
 250      255  
 10      AAT GCT AAA ATC ATA ATA GTC CAG TTG AAG GAA CCT GTA 816  
 Asn Ala Lys Ile Ile Ile Val Gln Leu Lys Glu Pro Val  
 260      265      270  
 15      GAA ATT AAT TGT ACA AGA CCC AGC AAC AAT ACA ATA AAA 855  
 Glu Ile Asn Cys Thr Arg Pro Ser Asn Asn Thr Ile Lys  
 275      280      285  
 20      GGT ATA CAC ATA GGA CCA GGG AGA GCA TTT TAT GCA ACA 894  
 Gly Ile His Ile Gly Pro Gly Arg Ala Phe Tyr Ala Thr  
 290      295  
 25      GGA GAC ATA CGA GGA GAT ATA AGA CAA GCA CAT TGT AAC 933  
 Gly Asp Ile Arg Gly Asp Ile Arg Gln Ala His Cys Asn  
 300      305      310  
 30      ATT AGT GGA GCA AAA TGG AAT AAC ACT TTA AAG AAG GTA 972  
 Ile Ser Gly Ala Lys Trp Asn Asn Thr Leu Lys Lys Val  
 315      320  
 35      GTT ATA AAA TTA AAA GAA CAA TTT CCA AAT AAA ACA ATA 1011  
 Val Ile Lys Leu Lys Glu Gln Phe Pro Asn Lys Thr Ile  
 325      330      335  
 40      GTC TTT AAC CAT TCC TCA GGA GGG GAC CCA GAA ATT GTA 1050  
 Val Phe Asn His Ser Ser Gly Gly Asp Pro Glu Ile Val  
 340      345      350  
 45      ATG CAC AGT TTT AAT TGT CAA GGG GAA TTT TTC TAC TGT 1089  
 Met His Ser Phe Asn Cys Gln Gly Glu Phe Phe Tyr Cys  
 355      360  
 50      AAT ACA ACG AAG CTG TTT AAT AGT ACT TGG AAT GAT ACT 1128  
 Asn Thr Thr Lys Leu Phe Asn Ser Thr Trp Asn Asp Thr  
 365      370      375  
 55      ACA GAG TCA AAT AAC AAT GAT AGT ACT ATT ACA CTC CCA 1167  
 Thr Glu Ser Asn Asn Asn Asp Ser Thr Ile Thr Leu Pro  
 380      385  
 60      TGC AGA ATA AAA CAA ATT ATA AAC ATG TGG CAG GAA GTA 1206  
 Cys Arg Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Val  
 390      395      400  
 65      GGA AAA GCA ATG TAT GCC CCT CCC ATC AGA GGA GAA ATT 1245  
 Gly Lys Ala Met Tyr Ala Pro Pro Ile Arg Gly Glu Ile  
 405      410      415  
 70      AAA TGT TCA TCA AAT ATT ACA GGA CTA CTG TTA ACA AGA 1284  
 Lys Cys Ser Ser Ile Thr Gly Leu Leu Leu Thr Arg  
 420      425  
 75      GAT GGT GGT ATT AAC ACT AGC GAT GCC ACC GAG ACC TTC 1323  
 Asp Gly Gly Ile Asn Thr Ser Asp Ala Thr Glu Thr Phe  
 430      435      440

AGA CCG GGA GGA GGA GAT ATG AGG GAC AAT TGG AGA AGT 1362  
 Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp Arg Ser  
 445 450

5 GAA TTA TAT AAA TAT AAA GTA GTG AAA ATT GAG CCA TTA 1401  
 Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile Glu Pro Leu  
 455 460 465

10 GGA GTA GCA CCC ACC AAG GCA AAG AGA AGA GTG GTG CAG 1440  
 Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln  
 470 475 480

AGA GAA AAA AGA GCA GTA ACA CTA GGA GCT ATG TTC CTT 1479  
 Arg Glu Lys Arg Ala Val Thr Leu Gly Ala Met Phe Leu  
 15 485 490

GGG TTC TTG GGA GCA TAA AGC TTC 1503  
 Gly Phe Leu Gly Ala Xaa Ser Phe  
 495 500 501

20 (2) INFORMATION FOR SEQ ID NO:4:  
 (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 501 amino acids  
 (B) TYPE: Amino Acid  
 25 (D) TOPOLOGY: Linear  
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:4:

Gly Val Pro Val Trp Lys Glu Ala Thr Thr Leu Phe Cys Ala  
 1 5 10 15

30 Ser Asp Ala Lys Ala Tyr Asp Thr Glu Val His Asn Val Trp Ala  
 20 25 30

Thr His Ala Cys Val Pro Thr Asp Pro Asn Pro Gln Glu Met Val  
 35 35 40 45

Leu Glu Asn Val Thr Glu Asp Phe Asn Met Trp Lys Asn Asp Met  
 50 55 60

40 Val Glu Cln Met His Glu Xaa Ile Ile Ser Leu Trp Asp Gln Ser  
 65 70 75

Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Ile Thr Leu Asn  
 80 85 90

45 Cys Thr Asn Trp Lys Glu Asn Asp Thr Lys Thr Asn Ser Ser Ser  
 95 100 105

50 Thr Thr Thr Asn Asn Ser Ser Ala Thr Ala Asn Ser Ser Thr  
 110 115 120

Thr Thr Asn Ser Ser Trp Gly Glu Ile Lys Glu Gly Glu Ile Lys  
 125 130 135

55 Asn Cys Ser Phe Asn Ile Thr Thr Gly Ile Arg Asp Lys Val Lys  
 140 145 150

Lys Glu Tyr Ala Leu Phe Tyr Ser Leu Asp Val Val Pro Ile Glu  
 155 160 165

60 Asn Asp Asn Thr Ser Tyr Arg Leu Arg Ser Cys Asn Thr Ser Val  
 170 175 180

65 Ile Thr Gln Ala Cys Pro Lys Val Thr Phe Glu Pro Ile Pro Ile  
 185 190 195

	His	Tyr	Cys	Thr	Pro	Ala	Gly	Phe	Ala	Ile	Leu	Lys	Cys	Lys	Asp
200															210
	Lys	Lys	Phe	Asn	Gly	Thr	Gly	Pro	Cys	Lys	Asn	Val	Ser	Thr	Val
5															225
	215									220					
	Gln	Cys	Thr	His	Gly	Ile	Lys	Pro	Val	Val	Ser	Thr	Gln	Leu	Leu
10															240
	230									235					
	Leu	Asn	Gly	Ser	Leu	Ala	Glu	Glu	Glu	Val	Ile	Ile	Arg	Ser	Ala
15															255
	245									250					
	Asn	Phe	Ser	Asn	Asn	Ala	Lys	Ile	Ile	Val	Gln	Leu	Lys	Glu	
20															270
	260									265					
	Pro	Val	Glu	Ile	Asn	Cys	Thr	Arg	Pro	Ser	Asn	Asn	Thr	Ile	Lys
25															285
	275									280					
	Gly	Ile	His	Ile	Gly	Pro	Gly	Arg	Ala	Phe	Tyr	Ala	Thr	Gly	Asp
30															300
	290									295					
	Ile	Arg	Gly	Asp	Ile	Arg	Gln	Ala	His	Cys	Asn	Ile	Ser	Gly	Ala
35															315
	305									310					
	Lys	Trp	Asn	Asn	Thr	Leu	Lys	Lys	Val	Val	Ile	Lys	Leu	Lys	Glu
40															330
	320									325					
	Gln	Phe	Pro	Asn	Lys	Thr	Ile	Val	Phe	Asn	His	Ser	Ser	Gly	Gly
45															345
	335									340					
	Asp	Pro	Glu	Ile	Val	Met	His	Ser	Phe	Asn	Cys	Gln	Gly	Glu	Phe
50															360
	350									355					
	Phe	Tyr	Cys	Asn	Thr	Thr	Lys	Leu	Phe	Asn	Ser	Thr	Trp	Asn	Asp
55															375
	365									370					
	Thr	Thr	Glu	Ser	Asn	Asn	Asn	Asp	Ser	Thr	Ile	Thr	Leu	Pro	Cys
60															390
	380									385					
	Arg	Ile	Lys	Gln	Ile	Ile	Asn	Met	Trp	Gln	Glu	Val	Gly	Lys	Ala
65															405
	395									400					
	Met	Tyr	Ala	Pro	Pro	Ile	Arg	Gly	Glu	Ile	Lys	Cys	Ser	Ser	Asn
70															420
	410									415					
	Ile	Thr	Gly	Leu	Leu	Leu	Thr	Arg	Asp	Gly	Gly	Ile	Asn	Thr	Ser
75															435
	425									430					
	Asp	Ala	Thr	Glu	Thr	Phe	Arg	Pro	Gly	Gly	Gly	Asp	Met	Arg	Asp
80															450
	440									445					
	Asn	Trp	Arg	Ser	Glu	Leu	Tyr	Lys	Tyr	Lys	Val	Val	Lys	Ile	Glu
85															465
	455									460					
	Pro	Leu	Gly	Val	Ala	Pro	Thr	Lys	Ala	Lys	Arg	Arg	Val	Val	Gln
90															480
	470									475					
	Arg	Glu	Lys	Arg	Ala	Val	Thr	Leu	Gly	Ala	Met	Phe	Leu	Gly	Phe
95															495
	485									490					
	Leu	Gly	Ala	Xaa	Ser	Phe									
100										500	501				

(2) INFORMATION FOR SEQ ID NO:5:  
 65 (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1461 base pairs

(B) TYPE: Nucleic Acid

(C) STRANDEDNESS: Single

(D) TOPOLOGY: Linear

5 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:5:

10	G	GTA	CCT	GTA	TGG	AAA	GAA	GCA	ACC	ACC	ACT	CTA	TTT	37
		Val	Pro	Val	Trp	Lys	Glu	Ala	Thr	Thr	Thr	Leu	Phe	
	1				5						10			
15	TGT	GCA	TCA	GAT	GCT	AAA	GCA	TAT	GAT	ACA	GAG	GTA	CAT	76
	Cys	Ala	Ser	Asp	Ala	Lys	Ala	Tyr	Asp	Thr	Glu	Val	His	
	15				20						25			
20	AAT	GTT	TGG	GCT	ACA	CAT	GCC	TGT	GTA	CCC	ACA	GAC	CCC	115
	Asn	Val	Trp	Ala	Thr	His	Ala	Cys	Val	Pro	Thr	Asp	Pro	
	30				35									
25	AAC	CCA	CAA	GAA	GTA	GTA	TTG	GAA	AAT	GTA	ACA	GAA	AAT	154
	Asn	Pro	Gln	Glu	Val	Val	Leu	Glu	Asn	Val	Thr	Glu	Asn	
	40				45					50				
30	TTT	AAC	ATG	TGG	AAA	aat	AAC	ATG	GTA	GAA	CAG	ATG	CAT	193
	Phe	Asn	Met	Trp	Lys	Asn	Asn	Met	Val	Glu	Gln	Met	His	
	55				60									
35	GAG	GAT	ATA	ATC	AGT	TTA	TGG	GAT	CAA	AGT	CTA	AAG	CCA	232
	Glu	Asp	Ile	Ile	Ser	Leu	Trp	Asp	Gln	Ser	Leu	Lys	Pro	
	65				70				75					
40	TGT	GTA	AAA	TTA	ACC	CCA	CTC	TGT	ACT	TTA	AAT	TGC	271	
	Cys	Val	Lys	Leu	Thr	Pro	Leu	Cys	Val	Thr	Leu	Asn	Cys	
	80				85				90					
45	ACT	AAT	TTG	GAG	AAT	GCT	AAT	ACC	GAG	AAT	GCT	AAT	310	
	Thr	Asn	Leu	Glu	Asn	Ala	Asn	Asn	Asn	Thr	Glu	Asn	Ala	
	95				100									
50	AAT	ACC	AAT	AAT	TAT	ACC	TTG	GGG	ATG	GAC	AGA	GGT	GAA	349
	Asn	Thr	Asn	Asn	Tyr	Thr	Leu	Gly	Met	Glu	Arg	Gly	Glu	
	105				110				115					
55	ATA	AAA	AAC	TGC	TCT	TTC	AAT	ATC	ACC	ACA	ACC	TTA	ACA	388
	Ile	Lys	Asn	Cys	Ser	Phe	Asn	Ile	Thr	Thr	Ser	Leu	Arg	
	120				125									
60	GAT	AAG	GTC	AAA	AAA	GAA	TAT	GCA	TTG	TTT	TAT	AAA	CTT	427
	Asp	Lys	Val	Lys	Lys	Glu	Tyr	Ala	Leu	Phe	Tyr	Lys	Leu	
	130				135				140					
65	GAT	GTA	CAA	ATA	GAT	AAT	AGT	ACC	AAC	TAT	AGG	CTG	466	
	Asp	Val	Val	Gln	Ile	Asp	Asn	Ser	Thr	Asn	Tyr	Arg	Leu	
	145				150				155					
70	ATA	AGT	TGT	AAT	ACC	TCA	GTC	ATT	ACA	CAG	GCC	TGT	CCA	505
	Ile	Ser	Cys	Asn	Thr	Ser	Val	Ile	Thr	Gln	Ala	Cys	Pro	
	160				165									
75	AAG	GTA	TCC	TTT	GAG	CTA	ATT	CCC	ATA	CAT	TAT	TGT	GCC	544
	Lys	Val	Ser	Phe	Glu	Leu	Ile	Pro	Ile	His	Tyr	Cys	Ala	
	170				175				180					
80	CCG	GCT	GGT	TTT	GCG	ATT	CTA	AAG	TGT	AAA	CAT	AAG	AAG	583
	Pro	Ala	Gly	Phe	Ala	Ile	Leu	Lys	Cys	Lys	Asp	Lys	Lys	
	185				190									

195.      TTC AAT GGA ACA GGA CCA TGT AAA AAT GTC AGC ACA GTA 622  
 Phe Asn Gly Thr Gly Pro Cys Lys Asn Val Ser Thr Val  
 200.      195.      205  
 5.      CAA TGT ACA CAT GGA ATT AGA CCA GTA GTA TCA ACT CAA 661  
 Gln Cys Thr His Gly Ile Arg Pro Val Val Ser Thr Gln  
 210.      210.      220  
 10.      CTA CTG TTA AAT GGC AGT CTA GCA GAA GAA GAG ATA GTA 700  
 Leu Leu Leu Asn Gly Ser Leu Ala Glu Glu Glu Ile Val  
 225.      225.      230  
 15.      ATT AGA TCT GAA AAT ATC ACA GAC AAT GCT AAA ACC ATA 739  
 Ile Arg Ser Glu Asn Ile Thr Asp Asn Ala Lys Thr Ile  
 235.      235.      240      245  
 20.      ATA GTG CAG CTA AAT GAA TCT ATA GTG ATT AAT TGT ACA 778  
 Ile Val Gln Leu Asn Glu Ser Ile Val Ile Asn Cys Thr  
 250.      250.      255  
 25.      AGA CCC AAT AAC AAC ACA AGA AAA AGT ATA AAT ATA GGA 817  
 Arg Pro Asn Asn Asn Thr Arg Lys Ser Ile Asn Ile Gly  
 260.      265.      270  
 30.      CCA GGG AGA GCA TTC TAT ACA ACA GGA GAC ATA ATA GGA 856  
 Pro Gly Arg Ala Phe Tyr Thr Thr Gly Asp Ile Ile Gly  
 275.      280.      285  
 35.      GAT ATA AGA CAA GCA CAT TGT AAC CTT AGT AAA ACA CAA 895  
 Asp Ile Arg Gln Ala His Cys Asn Leu Ser Lys Thr Gln  
 290.      290.      295  
 40.      TGG GAA AAA ACG TTA AGA CAG ATA GCT ATA AAA TTA GAA 934  
 Trp Glu Lys Thr Leu Arg Gln Ile Ala Ile Lys Leu Glu  
 300.      305.      310  
 45.      GAA AAA TTT AAG AAT AAA ACA ATA GCC TTT AAT AAA TCC 973  
 Glu Lys Phe Lys Asn Lys Thr Ile Ala Phe Asn Lys Ser  
 315.      315.      320  
 50.      TCA GGA GGG GAC CCA GAA ATT CTA ATG CAC AGT TTT AAT 1012  
 Ser Gly Gly Asp Pro Glu Ile Val Met His Ser Phe Asn  
 325.      330.      335  
 55.      TGT GGA GGG GAA TTT TTC TAC TGT AAT ACA ACA AAA CTG 1051  
 Cys Gly Gly Glu Phe Phe Tyr Cys Asn Thr Thr Lys Leu  
 340.      345.      350  
 60.      TTT AAT AGT ACC TGG AAT TTA ACA CAA CCG TTT AGT AAT 1090  
 Phe Asn Ser Thr Trp Asn Leu Thr Gln Pro Phe Ser Asn  
 355.      355.      360  
 65.      ACC GGG AAT CGT ACT GAA GAG TTA AAT ATT ACA CTC CCA 1129  
 Thr Gly Asn Arg Thr Glu Glu Leu Asn Ile Thr Leu Pro  
 365.      365.      370      375  
 70.      TGC AGA ATA AAA CAA ATC ATA AAC TTG TGG CAG GAA GTA 1168  
 Cys Arg Ile Lys Gln Ile Ile Asn Leu Trp Gln Glu Val  
 380.      380.      385  
 75.      GGC AAA GCA ATG TAT GCC CCT CCC ATC AGA GGA CAA ATT 1207  
 Gly Lys Ala Met Tyr Ala Pro Pro Ile Arg Gly Gln Ile  
 390.      395.      400

AGA TGT TCA TCA AAT ATT ACA GGG CTA CTA TTA ACA AGA 1246  
 Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg  
 405 410 415  
 5 GAT GGT GGA AGT AAC ACC GGT GAC AAC AGG ACT GAG ACC 1285  
 Asp Gly Gly Ser Asn Thr Gly Asp Asn Arg Thr Glu Thr  
 420 425  
 10 TTT AGA CCT GGA GGA GGA GAT ATG AGG GAC AAT TCG AGA 1324  
 Phe Arg Pro Gly Gly Asp Met Arg Asp Asn Trp Arg  
 430 435 440  
 15 AGT GAA TTA TAT AAA TAT AAA GTA CTA AGA ATT GAA CCA 1363  
 Ser Glu Leu Tyr Lys Tyr Lys Val Val Arg Ile Glu Pro  
 445 450  
 20 TTA GGA GTA GCA CCC ACC CAG GCA AAG AGA AGA GTG GTG 1402  
 Leu Gly Val Ala Pro Thr Gln Ala Lys Arg Arg Val Val  
 455 460 465  
 25 CAA AGA GAA AAA AGA GCA GTG GGG ATA GGA GCT ATG TTC 1441  
 Gln Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Met Phe  
 470 475 480  
 30 CTT GGG TTC TTG GGA GAT AA 1461  
 Leu Gly Phe Leu Gly Asp  
 485 486

(2) INFORMATION FOR SEQ ID NO:6:  
 30 (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 486 amino acids  
 (B) TYPE: Amino Acid  
 (D) TOPOLOGY: Linear  
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:6:  
 35 Val Pro Val Trp Lys Glu Ala Thr Thr Leu Phe Cys Ala Ser  
 1 5 10 15  
 Asp Ala Lys Ala Tyr Asp Thr Glu Val His Asn Val Trp Ala Thr  
 40 20 25 30  
 His Ala Cys Val Pro Thr Asp Pro Asn Pro Cln Glu Val Val Leu  
 35 40 45  
 45 Glu Asn Val Thr Glu Asn Phe Asn Met Trp Lys Asn Asn Met Val  
 50 55 60  
 Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp Asp Gln Ser Leu  
 65 70 75  
 50 Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu Asn Cys  
 80 85 90  
 Thr Asn Leu Glu Asn Ala Asn Asn Thr Glu Asn Ala Asn Asn Thr  
 55 95 100 105  
 Asn Asn Tyr Thr Leu Gly Met Glu Arg Gly Glu Ile Lys Asn Cys  
 110 115 120  
 60 Ser Phe Asn Ile Thr Thr Ser Leu Arg Asp Lys Val Lys Lys Glu  
 125 130 135  
 Tyr Ala Leu Phe Tyr Lys Leu Asp Val Val Cln Ile Asp Asn Ser  
 140 145 150  
 65

	Thr Asn Tyr Arg Leu Ile Ser Cys Asn Thr Ser Val Ile Thr Gln		
	155	160	165
5	Ala Cys Pro Lys Val Ser Phe Glu Leu Ile Pro Ile His Tyr Cys		
	170	175	180
	Ala Pro Ala Gly Phe Ala Ile Leu Lys Cys Lys Asp Lys Lys Phe		
	185	190	195
10	Asn Gly Thr Gly Pro Cys Lys Asn Val Ser Thr Val Gln Cys Thr		
	200	205	210
	His Gly Ile Arg Pro Val Val Ser Thr Gln Leu Leu Leu Asn Gly		
	215	220	225
15	Ser Leu Ala Glu Glu Glu Ile Val Ile Arg Ser Glu Asn Ile Thr		
	230	235	240
20	Asp Asn Ala Lys Thr Ile Ile Val Gln Leu Asn Glu Ser Ile Val		
	245	250	255
	Ile Asn Cys Thr Arg Pro Asn Asn Asn Thr Arg Lys Ser Ile Asn		
	260	265	270
25	Ile Gly Pro Gly Arg Ala Phe Tyr Thr Thr Gly Asp Ile Ile Gly		
	275	280	285
	Asp Ile Arg Gln Ala His Cys Asn Leu Ser Lys Thr Gln Trp Glu		
	290	295	300
30	Lys Thr Leu Arg Gln Ile Ala Ile Lys Leu Glu Glu Lys Phe Lys		
	305	310	315
35	Asn Lys Thr Ile Ala Phe Asn Lys Ser Ser Gly Gly Asp Pro Glu		
	320	325	330
	Ile Val Met His Ser Phe Asn Cys Gly Gly Glu Phe Phe Tyr Cys		
	335	340	345
40	Asn Thr Thr Lys Leu Phe Asn Ser Thr Trp Asn Leu Thr Gln Pro		
	350	355	360
	Phe Ser Asn Thr Gly Asn Arg Thr Glu Glu Leu Asn Ile Thr Leu		
	365	370	375
45	Pro Cys Arg Ile Lys Gln Ile Ile Asn Leu Trp Gln Glu Val Gly		
	380	385	390
50	Lys Ala Met Tyr Ala Pro Pro Ile Arg Gly Gln Ile Arg Cys Ser		
	395	400	405
	Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly Ser Asn		
	410	415	420
55	Thr Gly Asp Asn Arg Thr Glu Thr Phe Arg Pro Gly Gly Asp		
	425	430	435
	Met Arg Asp Asn Trp Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val		
	440	445	450
60	Arg Ile Glu Pro Leu Gly Val Ala Pro Thr Gln Ala Lys Arg Arg		
	455	460	465
65	Val Val Gln Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Met Phe		
	470	475	480

Leu Gly Phe Leu Gly Asp  
485 486

## (2) INFORMATION FOR SEQ ID NO:7:

## 5 (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 1474 base pairs
- (B) TYPE: Nucleic Acid
- (C) STRANDEDNESS: Single
- (D) TOPOLOGY: Linear

## 10 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:7:

G GCA CCT GTG TGG AAA GAA GCA ACC ACC ACT CTA TTT 37  
Val Pro Val Trp Lys Glu Ala Thr Thr Thr Leu Phe  
15 1 5 10

TGT GCA TCA GAT GCT AAA GCA TAT GAT ACA GAG GTC CAT 76  
Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu Val His  
20 15 20 25

AAT GTT TGG GCT ACA CAT GCC TGT GTC CCC ACA GAC CCC 115  
Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro  
25 30 35

AAC CCA CAA GAA GTC GTC TTG GAA AAT GTC ACA GAA AAT 154  
Asn Pro Gln Glu Val Val Leu Glu Asn Val Thr Glu Asn  
30 40 45 50

TTT AAC ATG TGG AAA AAT AAC ATG GTC GAA CAG ATG CAT 193  
Phe Asn Met Trp Lys Asn Asn Met Val Glu Gln Met His  
35 55 60

GAG GAT ATA ATC AGT TTA TGG GAT CAA AGT CTC AAG CCA 232  
Glu Asp Ile Ile Ser Leu Trp Asp Gln Ser Leu Lys Pro  
40 65 70 75

TGT GTC AAA TTA ACC CCA CTC TGT GTT ACT TTA AAT TGC 271  
Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu Asn Cys  
45 80 85 90

ACT AAT TTG GAG AAT GCT AAT AAT ACC GAG AAT GCT AAT 310  
Thr Asn Leu Glu Asn Ala Asn Asn Thr Glu Asn Ala Asn  
50 95 100

AAT ACC AAT AAT TAT ACC TTG GGG ATG GAG AGA GGT GAA 349  
Asn Thr Asn Asn Tyr Thr Leu Gly Met Glu Arg Gly Glu  
55 105 110 115

AGA AAA AAC TGC TCT TTC AAT ATC ACC ACA AGC TTA AGA 388  
Arg Lys Asn Cys Ser Phe Asn Ile Thr Thr Ser Leu Arg  
60 120 125

GAT AAG GGG AAA AAA GAA TAT GCA TTG TTT TAT AAA CTT 427  
Asp Lys Gly Lys Lys Glu Tyr Ala Leu Phe Tyr Lys Leu  
65 130 135 140

GAT GTC GTC CAA ATA GAT AAT AGT ACC AAC TAT AGG CTG 466  
Asp Val Val Gln Ile Asp Asn Ser Thr Asn Tyr Arg Leu  
70 145 150 155

ATA AGT TGT AAT ACC TCA GTC ATT ACA CAC GCC TGT CCA 505  
Ile Ser Cys Asn Thr Ser Val Ile Thr Gln Ala Cys Pro  
75 160 165

AAG GTA TCC TTT GAG CCA ATT CCC ATA CAT TAT TGT GCC 544  
 Lys Val Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala  
 170 175 180  
 5 CCG GCT GGT TTT GCG ATT CTA AAG TGT AAA GAT AAG AAG 583  
 Pro Ala Gly Phe Ala Ile Leu Lys Cys Lys Asp Lys Lys  
 185 190  
 10 TTC AAT CGA ACA GGA CCA TGT AAA AAT GTC AGG ACA GTA 622  
 Phe Asn Gly Thr Gly Pro Cys Lys Asn Val Arg Thr Val  
 195 200 205  
 15 CAA TGT ACA CAT GGA ATT AGA CCA GTC GTC TCA ACT CAA 661  
 Gln Cys Thr His Gly Ile Arg Pro Val Val Ser Thr Gln  
 210 215 220  
 20 CTA CTG TTA AAT GGC AGT CTA GCA GAA GAA GAG ATA GTA 700  
 Leu Leu Leu Asn Gly Ser Leu Ala Glu Glu Glu Ile Val  
 225 230  
 25 ATT AGA TCT GAA AAT ATC ACA GAC AAT GCT AAA ACC ATA 739  
 Ile Arg Ser Glu Asn Ile Thr Asp Asn Ala Lys Thr Ile  
 235 240 245  
 30 ATA CTG CAG CTA AAT GAA TCT ATA GTG ATT AAT TGT ACA 778  
 Ile Val Gln Leu Asn Glu Ser Ile Val Ile Asn Cys Thr  
 250 255  
 35 AGA CCC AAT AAC AAC ACA AGA AAA AGT ATA AAT ATA GGA 817  
 Arg Pro Asn Asn Asn Thr Arg Lys Ser Ile Asn Ile Gly  
 260 265 270  
 40 CCA GGG AGA GCA TTC TAT ACA ACA GGA GAC ATA ATA GGA 856  
 Pro Gly Arg Ala Phe Tyr Thr Thr Gly Asp Ile Ile Gly  
 275 280 285  
 45 GAT ATA AGA CAA GCA CAT TGT AAC CTT AGT AAA ACA CAA 895  
 Asp Ile Arg Gln Ala His Cys Asn Leu Ser Lys Thr Gln  
 290 295  
 50 TGG GAA AAA ACG TTA AGA CAG ATA GCT ATA AAA TTA GAA 934  
 Trp Glu Lys Thr Leu Arg Gln Ile Ala Ile Lys Leu Glu  
 300 305 310  
 55 GAA AAA TTT AAG AAT AAA ACA ATA GCC TTT AAT AAA TCC 973  
 Glu Lys Phe Lys Asn Lys Thr Ile Ala Phe Asn Lys Ser  
 315 320  
 60 TCA GGA GGG GAC CCA GAA ATT GTA ATG CAC AGT TTT AAT 1012  
 Ser Gly Gly Asp Pro Glu Ile Val Met His Ser Phe Asn  
 325 330 335  
 65 TGT GGA GGG GGA TTT TTC TAC TGT AGT ACG AGA AAA CTG 1051  
 Cys Gly Gly Gly Phe Phe Tyr Cys Ser Thr Arg Lys Leu  
 340 345 350  
 70 TTT AAT AGT ACC TGG AAT TTA ACA CAA CCG TTT AGT AAT 1090  
 Phe Asn Ser Thr Trp Asn Leu Thr Gln Pro Phe Ser Asn  
 355 360  
 75 ACC GGG GAT CGT ACT GAA GAG TTA AAT ATT ACA CTC CCA 1129  
 Thr Gly Asp Arg Thr Glu Glu Leu Asn Ile Thr Leu Pro  
 365 370 375

TGC AGA ATA AAA CAA ATC ATA AAC TTG TGG CAG GAA GTA 1168  
 Cys Arg Ile Lys Gln Ile Ile Asn Leu Trp Gln Glu Val  
 380 385  
 5 GGC AAA GCA ATG TAT GCC CCT CCC ATC AGA GGA CAA ATT 1207  
 Gly Lys Ala Met Tyr Ala Pro Pro Ile Arg Gly Gln Ile  
 390 395 400  
 10 AGA TGT TCA TCA AAT ATT ACA GGG CTA CTA TTA AGG AGA 1246  
 Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Arg Arg  
 405 410 415  
 15 GAT GGT GGA AGT AAC ACC AGT GAC AAC CAG ACT GAG ACC 1285  
 Asp Gly Gly Ser Asn Thr Ser Asp Asn Gln Thr Glu Thr  
 420 425  
 20 TTT AGA CCT GGG GGA GGA GAT ATG AGG GAC AAG TGG AGA 1324  
 Phe Arg Pro Gly Gly Asp Met Arg Asp Lys Trp Arg  
 430 435 440  
 25 AGT GAA TTA TAT AAA TAT AAA GTA GTA AGA ATT GAA CCA 1363  
 Ser Glu Leu Tyr Lys Tyr Lys Val Val Arg Ile Glu Pro  
 445 450  
 30 TTA GGA GTA GCA CCC ACC CAG GCA AAG AGA AGA CTG CTG 1402  
 Leu Gly Val Ala Pro Thr Gln Ala Lys Arg Arg Val Val  
 455 460 465  
 35 CAA AGA GAA AAA AGA GCA GTG GGG ATA GGA CCT ATC TTC 1441  
 Gln Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Met Phe  
 470 475 480  
 CTT AGG TTC TTA GGA GAT AAA GCT TCT AGA GTC 1474  
 Leu Arg Phe Leu Gly Asp Lys Ala Ser Arg Val  
 485 490 491

## (2) INFORMATION FOR SEQ ID NO:8:

## (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 491 amino acids

40 (B) TYPE: Amino Acid

(D) TOPOLOGY: Linear

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:8:

Val Pro Val Trp Lys Glu Ala Thr Thr Thr Leu Phe Cys Ala Ser  
 45 1 5 10 15  
 Asp Ala Lys Ala Tyr Asp Thr Glu Val His Asn Val Trp Ala Thr  
 20 25 30  
 50 His Ala Cys Val Pro Thr Asp Pro Asn Pro Gln Glu Val Val Leu  
 35 40 45  
 Glu Asn Val Thr Glu Asn Phe Asn Met Trp Lys Asn Asn Met Val  
 55 50 55 60  
 55 Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp Asp Gln Ser Leu  
 65 70 75  
 60 Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu Asn Cys  
 80 85 90  
 Thr Asn Leu Glu Asn Ala Asn Asn Thr Glu Asn Ala Asn Asn Thr  
 95 100 105

	Asn Asn Tyr Thr Leu Gly Met Glu Arg Gly Glu Arg Lys Asn Cys	
	110 115 120	
5	Ser Phe Asn Ile Thr Thr Ser Leu Arg Asp Lys Gly Lys Lys Glu	
	125 130 135	
	Tyr Ala Leu Phe Tyr Lys Leu Asp Val Val Gln Ile Asp Asn Ser	
	140 145 150	
10	Thr Asn Tyr Arg Leu Ile Ser Cys Asn Thr Ser Val Ile Thr Gln	
	155 160 165	
	Ala Cys Pro Lys Val Ser Phe Glu Pro Ile Pro Ile His Tyr Cys	
	170 175 180	
15	Ala Pro Ala Gly Phe Ala Ile Leu Lys Cys Lys Asp Lys Lys Phe	
	185 190 195	
20	Asn Gly Thr Gly Pro Cys Lys Asn Val Arg Thr Val Gln Cys Thr	
	200 205 210	
	His Gly Ile Arg Pro Val Val Ser Thr Gln Leu Leu Leu Asn Gly	
	215 220 225	
25	Ser Leu Ala Glu Glu Glu Ile Val Ile Arg Ser Glu Asn Ile Thr	
	230 235 240	
	Asp Asn Ala Lys Thr Ile Ile Val Gln Leu Asn Glu Ser Ile Val	
	245 250 255	
30	Ile Asn Cys Thr Arg Pro Asn Asn Asn Thr Arg Lys Ser Ile Asn	
	260 265 270	
	Ile Gly Pro Gly Arg Ala Phe Tyr Thr Thr Gly Asp Ile Ile Gly	
35	275 280 285	
	Asp Ile Arg Gln Ala His Cys Asn Leu Ser Lys Thr Gln Trp Glu	
	290 295 300	
40	Lys Thr Leu Arg Gln Ile Ala Ile Lys Leu Glu Glu Lys Phe Lys	
	305 310 315	
	Asn Lys Thr Ile Ala Phe Asn Lys Ser Ser Gly Gly Asp Pro Glu	
	320 325 330	
45	Ile Val Met His Ser Phe Asn Cys Gly Gly Phe Phe Tyr Cys	
	335 340 345	
50	Ser Thr Arg Lys Leu Phe Asn Ser Thr Trp Asn Leu Thr Gln Pro	
	350 355 360	
	Phe Ser Asn Thr Gly Asp Arg Thr Glu Glu Leu Asn Ile Thr Leu	
	365 370 375	
55	Pro Cys Arg Ile Lys Gln Ile Ile Asn Leu Trp Gln Glu Val Gly	
	380 385 390	
	Lys Ala Met Tyr Ala Pro Pro Ile Arg Gly Gln Ile Arg Cys Ser	
	395 400 405	
60	Ser Asn Ile Thr Gly Leu Leu Leu Arg Arg Asp Gly Gly Ser Asn	
	410 415 420	
	Thr Ser Asp Asn Gln Thr Glu Thr Phe Arg Pro Gly Gly Asp	
65	425 430 435	

	Met Arg Asp Lys Trp Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val		
	440	445	450
5	Arg Ile Glu Pro Leu Gly Val Ala Pro Thr Gln Ala Lys Arg Arg		
	455	460	465
	Val Val Gln Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Met Phe		
	470	475	480
10	Leu Arg Phe Leu Gly Asp Lys Ala Ser Arg Val		
	485	490	491

## (2). INFORMATION FOR SEQ ID NO:9:

## (i) SEQUENCE CHARACTERISTICS:

15 (A) LENGTH: 1512 base pairs  
 (B) TYPE: Nucleic Acid  
 (C) STRANDEDNESS: Single  
 (D) TOPOLOGY: Linear

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:9:

20	CTC GAG GTA CCT GTA TGG AAA GAA GCA ACT ACC ACT 36		
	Leu Glu Val Pro Val Trp Lys Glu Ala Thr Thr Thr		
	1 5 10		
25	CTA TTT TGT GCA TCA GAT GCT AAA GCA TAT AAT ACA GAG 75		
	Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asn Thr Glu		
	15 20 25		
30	AAA CAT AAT GTT TGG GCC ACA CAC GCC TGT GTA CCC ACA 114		
	Lys His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr		
	30 35		
35	GAT CCC AAC CCA CAA GAA GTA GTA TTG GGA AAT GTG ACA 153		
	Asp Pro Asn Pro Gln Glu Val Val Leu Gly Asn Val Thr		
	40 45 50		
40	GAA AAT TTT AAC ATG TGG AAA AAT AAC ATG GTA GAA CAA 192		
	Glu Asn Phe Asn Met Trp Lys Asn Asn Met Val Glu Gln		
	55 60		
45	ATG CAT GAA GAT ATA ATC AGT TTA TGG GAT CAA AGT CTA 231		
	Met His Glu Asp Ile Ile Ser Leu Trp Asp Gln Ser Leu		
	65 70 75		
50	AAG CCA TGT GTA AAA TTA ACC CCA CTC TCT GTT ACT TTA 270		
	Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu		
	80 85 90		
55	AAT TGC ACT GAT GAT TTA GGG AAT GCT ACT AAT ACC AAT 309		
	Asn Cys Thr Asp Asp Leu Gly Asn Ala Thr Asn Thr Asn		
	95 100		
60	AGT AGT GCC ACT ACC AAT AGT AGT AGT TGG GAA GAA ATG 348		
	Ser Ser Ala Thr Thr Asn Ser Ser Trp Glu Glu Met		
	105 110 115		
65	AAG GGG GAA ATG AAA AGA TGC TCT TTC AAT ATC ACC ACA 387		
	Lys Gly Glu Met Lys Arg Cys Ser Phe Asn Ile Thr Thr		
	120 125		
70	AGC ATA AGA GAT AAG ATT AAG AAA CAA CAT GCA CTT TTC 426		
	Ser Ile Arg Asp Lys Ile Lys Lys Glu His Ala Leu Phe		
	130 135 140		

TAT AGA CTT GAT GTA GTA CCA ATA GAT AAT GAT AAT ACC 465  
 Tyr Arg Leu Asp Val Val Pro Ile Asp Asn Asp Asn Thr  
 145 150 155  
 5 ACA TAT AGG TTG ATA AAT TGT AAT ACC TCA GTC ATT ACA 504  
 Thr Tyr Arg Leu Ile Asn Cys Asn Thr Ser Val Ile Thr  
 160 165  
 10 CAG GCC TGT CCA AAG GTA TCA TTT GAG CCA ATT CCC ATA 543  
 Gln Ala Cys Pro Lys Val Ser Phe Glu Pro Ile Pro Ile  
 170 175 180  
 15 CAT TTT TGT GCC CCG GCT GGT TTT GCG ATT CTA AAG TGT 582  
 His Phe Cys Ala Pro Ala Gly Phe Ala Ile Leu Lys Cys  
 185 190  
 20 AAT AAT AAG ACG TTC GAG CCA AAA GGA CCA TGT AAA AAT 621  
 Asn Asn Lys Thr Phe Glu Gly Lys Gly Pro Cys Lys Asn  
 195 200 205  
 25 GTC AGT ACA GTA CAA TGC ACA CAT GGA ATT AGG CCA GTA 660  
 Val Ser Thr Val Gln Cys Thr His Gly Ile Arg Pro Val  
 210 215 220  
 30 GTG TCA ACT CAA CTG CTG TTA AAT GGC ACT CTA GCA GAA 699  
 Val Ser Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu  
 225 230  
 35 GAA GAG GTA ATA ATT AGA TCT GAC AAT ATC ACA GAC AAT 738  
 Glu Glu Val Ile Ile Arg Ser Asp Asn Ile Thr Asp Asn  
 235 240 245  
 40 ACT AAA ACC ATT ATA GTA CAG CTA AAC GAA TCT CTA CTA 777  
 Thr Lys Thr Ile Ile Val Gln Leu Asn Glu Ser Val Val  
 250 255  
 45 ATT AAT TGT ACA AGA CCC AAC AAC AAT ACA AGA AAA AGT 816  
 Ile Asn Cys Thr Arg Pro Asn Asn Asn Thr Arg Lys Ser  
 260 265 270  
 50 ATA CAT ATA GGA CCA GGG AGT GCA TTT TTT GCA ACA GGA 855  
 Ile His Ile Gly Pro Gly Ser Ala Phe Phe Ala Thr Gly  
 275 280 285  
 55 GAA ATA ATA GGA GAT ATA AGA CAA GCA CAC TGT AAC CTT 894  
 Glu Ile Ile Gly Asp Ile Arg Gln Ala His Cys Asn Leu  
 290 295  
 60 AGT AGA ACA CAA TGG AAT AAC ACT TTA GGA AAG ATA GTC 933  
 Ser Arg Thr Gln Trp Asn Asn Thr Leu Gly Lys Ile Val  
 300 305 310  
 65 ATA AAA TTA AGA GAA CAA TTT AGA AAA CAA TTT GGA GAA 972  
 Ile Lys Leu Arg Glu Gln Phe Arg Lys Gln Phe Gly Glu  
 315 320  
 70 AAA ACA ATA GTC TTT AAT CGA TCC TCA GGA GGG GAC CCG 1011  
 Lys Thr Ile Val Phe Asn Arg Ser Ser Gly Gly Asp Pro  
 325 330 335  
 75 GAA ATT GCA ATG CAC AGT TTT AAT TGT GGA GGG GAA TTT 1050  
 Glu Ile Ala Met His Ser Phe Asn Cys Gly Gly Glu Phe  
 340 345 350

TTC TAC TGT AAC ACA ACA GCA CTG TTT AAT AGT ACC TGG 1089  
 Phe Tyr Cys Asn Thr Thr Ala Leu Phe Asn Ser Thr Trp  
 355 360

5 AAT GTT ACT AAA GGG TTG AAT AAC ACT GAA GGA AAT AGC 1128  
 Asn Val Thr Lys Gly Leu Asn Asn Thr Glu Gly Asn Ser  
 365 370 375

10 ACA GGA GAT GAA AAT ATC ATA CTC CCA TGT AGA ATA AAA 1167  
 Thr Gly Asp Glu Asn Ile Ile Leu Pro Cys Arg Ile Lys  
 380 385

15 CAA ATT ATA AAC ATG TGG CAG GAA GTC GGA AAA GCA ATG 1206  
 Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys Ala Met  
 390 395 400

20 TAT GCC CCT CCC ATC AGT GGA CAA ATT AGA TGT TCA TCA 1245  
 Tyr Ala Pro Pro Ile Ser Gly Gln Ile Arg Cys Ser Ser  
 405 410 415

25 AAC ATT ACA GGG CTG CTA CTA ACA AGA GAT GGT GGT AGT 1284  
 Asn Ile Thr Gly Leu Leu Thr Arg Asp Gly Gly Ser  
 420 425

30 AAG AAC GAG AGC ATC ACC ACC GAG GTC TTC AGA CCT GGA 1323  
 Lys Asn Glu Ser Ile Thr Thr Glu Val Phe Arg Pro Gly  
 430 435 440

35 GGA GGA GAT ATG AGG GAC AAT TGG ACA ACT CAA TTA TAT 1362  
 Gly Gly Asp Met Arg Asp Asn Trp Arg Ser Glu Leu Tyr  
 445 450

40 AAA TAT AAA GTA GTA AAA ATT GAA CCA TTA GGA GTC GCG 1401  
 Lys Tyr Lys Val Val Lys Ile Glu Pro Leu Gly Val Ala  
 455 460 465

45 CCC ACC AAC GCA AAG AGA AGA GTG GTG CAG AGA GAA AAA 1440  
 Pro Thr Lys Ala Lys Arg Arg Val Val Gln Arg Glu Lys  
 470 475 480

50 AGA GCA GTG GGA ACA ATA GGA GCT ATG TTC CTT GGG TTC 1479  
 Arg Ala Val Gly Thr Ile Gly Ala Met Phe Leu Gly Phe  
 485 490

55 TTG GGA GCA TAA AGC TTC TAG AGT CGA CCT GCA 1512  
 Leu Gly Ala Xaa Ser Phe Xaa Ser Arg Pro Ala  
 495 500 504

(2) INFORMATION FOR SEQ ID NO:10:  
 50 (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 504 amino acids  
 (B) TYPE: Amino Acid  
 (D) TOPOLOGY: Linear  
 55 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:10:  
 Leu Glu Val Pro Val Trp Lys Glu Ala Thr Thr Thr Leu Phe Cys  
 1 5 10 15  
 Ala Ser Asp Ala Lys Ala Tyr Asn Thr Glu Lys His Asn Val Trp  
 60 20 25 30  
 Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn Pro Gln Glu Val  
 35 40 45

Val Leu Gly Asn Val Thr Glu Asn Phe Asn Met Trp Lys Asn Asn  
50 55 60

Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp Asp Gln  
5 65 70 75

Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu  
80 85 90

10 Asn Cys Thr Asp Asp Leu Gly Asn Ala Thr Asn Thr Asn Ser Ser  
95 100 105

Ala Thr Thr Asn Ser Ser Trp Glu Glu Met Lys Gly Glu Met  
110 115 120

15 Lys Arg Cys Ser Phe Asn Ile Thr Thr Ser Ile Arg Asp Lys Ile  
125 130 135

Lys Lys Glu His Ala Leu Phe Tyr Arg Leu Asp Val Val Pro Ile  
20 140 145 150

Asp Asn Asp Asn Thr Thr Tyr Arg Leu Ile Asn Cys Asn Thr Ser  
155 160 165

25 Val Ile Thr Gln Ala Cys Pro Lys Val Ser Phe Glu Pro Ile Pro  
170 175 180

Ile His Phe Cys Ala Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn  
185 190 195

30 Asn Lys Thr Phe Glu Gly Lys Gly Pro Cys Lys Asn Val Ser Thr  
200 205 210

Val Gln Cys Thr His Gly Ile Arg Pro Val Val Ser Thr Gln Leu  
35 215 220 225

Leu Leu Asn Gly Ser Leu Ala Glu Glu Val Ile Ile Arg Ser  
230 235 240

40 Asp Asn Ile Thr Asp Asn Thr Lys Thr Ile Ile Val Gln Leu Asn  
245 250 255

Glu Ser Val Val Ile Asn Cys Thr Arg Pro Asn Asn Asn Thr Arg  
260 265 270

45 Lys Ser Ile His Ile Gly Pro Gly Ser Ala Phe Phe Ala Thr Gly  
275 280 285

Glu Ile Ile Gly Asp Ile Arg Gln Ala His Cys Asn Leu Ser Arg  
50 290 295 300

Thr Gln Trp Asn Asn Thr Leu Gly Lys Ile Val Ile Lys Leu Arg  
305 310 315

55 Glu Gln Phe Arg Lys Gln Phe Gly Glu Lys Thr Ile Val Phe Asn  
320 325 330

Arg Ser Ser Gly Gly Asp Pro Glu Ile Ala Met His Ser Phe Asn  
335 340 345

60 Cys Gly Gly Glu Phe Phe Tyr Cys Asn Thr Thr Ala Leu Phe Asn  
350 355 360

Ser Thr Trp Asn Val Thr Lys Gly Leu Asn Asn Thr Glu Gly Asn  
65 365 370 375

Ser Thr Gly Asp Glu Asn Ile Ile Leu Pro Cys Arg Ile Lys Gln  
 380 385 390  
 Ile Ile Asn Met Trp Gln Glu Val Gly Lys Ala Met Tyr Ala Pro  
 5 395 400 405  
 Pro Ile Ser Gly Gln Ile Arg Cys Ser Ser Asn Ile Thr Gly Leu  
 410 415 420  
 10 Leu Leu Thr Arg Asp Gly Gly Ser Lys Asn Glu Ser Ile Thr Thr  
 425 430 435  
 Glu Val Phe Arg Pro Gly Gly Asp Met Arg Asp Asn Trp Arg  
 15 440 445 450  
 Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile Glu Pro Leu Gly  
 455 460 465  
 Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln Arg Glu Lys  
 20 470 475 480  
 Arg Ala Val Gly Thr Ile Gly Ala Met Phe Leu Gly Phe Leu Gly  
 485 490 495  
 25 Ala Xaa Ser Phe Xaa Ser Arg Pro Ala  
 500 504

## (2) INFORMATION FOR SEQ ID NO:11:

## (i) SEQUENCE CHARACTERISTICS:

30 (A) LENGTH: 1501 base pairs  
 (B) TYPE: Nucleic Acid  
 (C) STRANDEDNESS: Single  
 (D) TOPOLOGY: Linear

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:11:

35 CTC GAG GTA CCT GTG TGG AAA GAA GCA ACT ACC ACT 36  
 Leu Glu Val Pro Val Trp Lys Glu Ala Thr Thr Thr  
 1 5 10

40 CTA TTT TGT GCA TCA GAT GCT AAA GCA TAT AAT ACA GAG 75  
 Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asn Thr Glu  
 15 20 25

45 AAA CAT AAT GTT TGG GCC ACA CAC GCC TGT GTA CCC ACA 114  
 Lys His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr  
 30 35

50 GAT CCC AAC CCA CAA GAA GTA GTA TTG GGA AAT GTG ACA 153  
 Asp Pro Asn Pro Gln Glu Val Val Leu Gly Asn Val Thr  
 40 45 50

55 GAA AAT TTT AAC ATG TGG AAA AAT AAC ATG GTA GAA CAA 192  
 Glu Asn Phe Asn Met Trp Lys Asn Asn Met Val Glu Gln  
 55 60

60 ATG CAT GAA GAT ATA ATC ACT TTA TGG GAT CAA AGT CTA 231  
 Met His Glu Asp Ile Ile Ser Leu Trp Asp Gln Ser Leu  
 65 70 75

65 AAG CCA TGT GTA AAA TTA ACC CCA CTC TGT CTT ACT TTA 270  
 Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu  
 80 85 90

AAT TGC ACT GAT GAT TTA GGG AAT GCT ACT AAT ACC AAT 309  
 Asn Cys Thr Asp Asp Leu Gly Asn Ala Thr Asn Thr Asn  
 95 100  
 5 AGC AGT CCC ACT ACC AAT AGT AGT TGG CAA GAA ATG 348  
 Ser Ser Ala Thr Thr Asn Ser Ser Trp Glu Glu Met  
 105 110 115  
 10 AAG GGG GAA ATG AAA AGG TGC TCT TTC AAT ATC ACC ACA 387  
 Lys Gly Glu Met Lys Arg Cys Ser Phe Asn Ile Thr Thr  
 120 125  
 15 AGC ATA AGA GAT AAG ATT AAG AAA GAA CAT GCA CTT TTC 426  
 Ser Ile Arg Asp Lys Ile Lys Lys Glu His Ala Leu Phe  
 130 135 140  
 20 TAT AGA CTT GAT GTA CCA ATA GAT AAT GAT AAT ACC 465  
 Tyr Arg Leu Asp Val Val Pro Ile Asp Asn Asp Asn Thr  
 145 150 155  
 25 ACA TAT AGG TTG ATA AAT TGT AAT ACC TCA GTC ATT ACA 504  
 Thr Tyr Arg Leu Ile Asn Cys Asn Thr Ser Val Ile Thr  
 160 165  
 30 CAG GCC TGT CCA AAG GTA TCA TTT GAG CCA ATT CCC ATA 543  
 Gln Ala Cys Pro Lys Val Ser Phe Glu Pro Ile Pro Ile  
 170 175 180  
 35 CAT TTT TGT GCC CCG GCT GGT TTT GCG ATT CTA AAG TGT 582  
 His Phe Cys Ala Pro Ala Gly Phe Ala Ile Leu Lys Cys  
 185 190  
 40 AAT AAT AAG ACG TTC GAG GGA AAA GGA CCA TGT AAA AAT 621  
 Asn Asn Lys Thr Phe Glu Gly Lys Gly Pro Cys Lys Asn  
 195 200 205  
 45 GTC AGT ACA GTA CAA TGC ACA CAT GGA ATT AGG CCA GTA 660  
 Val Ser Thr Val Gln Cys Thr His Gly Ile Arg Pro Val  
 210 215 220  
 50 GTG TCA ACT CAA CTG CTG TTA AAT GGC AGT CTA GCA GAA 699  
 Val Ser Thr Gln Leu Leu Asn Gly Ser Leu Ala Glu  
 225 230  
 55 GAA GAG GTA ATA ATT AGA TCT GGC AAT ATC ACA GAC AAT 738  
 Glu Glu Val Ile Ile Arg Ser Gly Asn Ile Thr Asp Asn  
 235 240 245  
 60 ACT AAA ACC ATT ATA GTA CAG CTA AAC GAA TCT GTA GTA 777  
 Thr Lys Thr Ile Ile Val Gln Leu Asn Glu Ser Val Val  
 250 255  
 65 ATT AAT TGT ACA AGA TCC AAC AAC AAT ACA AGA AAA AGT 816  
 Ile Asn Cys Thr Arg Ser Asn Asn Asn Thr Arg Lys Ser  
 260 265 270  
 70 ATA CAT ATA GGA CCA GGG AGT GCA TTT TTT GCA ACA GGA 855  
 Ile His Ile Gly Pro Gly Ser Ala Phe Phe Ala Thr Gly  
 275 280 285  
 75 GAA ATA ATA GGA GAT ATA AGA CAA GCA CAC TGT AAC CTT 894  
 Glu Ile Ile Gly Asp Ile Arg Gln Ala His Cys Asn Leu  
 290 295

AGT AGA ACA CAA TGG AAT AAC ACT TTA GGA AAG ATA GTC 933  
 Ser Arg Thr Gln Trp Asn Asn Thr Leu Gly Lys Ile Val  
 300 305 310  
 5 ATA AAA TTA AGA GAA CAA TTT AGA AAA CAN TTT GGA GAA 972  
 Ile Lys Leu Arg Glu Gln Phe Arg Lys Gln Phe Gly Glu  
 315 320  
 10 AAA ACA ATA GTC TTT AAT CGA TCC TCA GGA GGG GAC CCG 1011  
 Lys Thr Ile Val Phe Asn Arg Ser Ser Gly Gly Asp Pro  
 325 330 335  
 15 GAA ATT GCA ATG CAC AGT TTT AAT TGT GGA GGG GAA TTT 1050  
 Glu Ile Ala Met His Ser Phe Asn Cys Gly Gly Glu Phe  
 340 345 350  
 20 TTC TAC TGT AAC ACA ACA GCA CTG TTT AAT AGT ACC TGG 1089  
 Phe Tyr Cys Asn Thr Thr Ala Leu Phe Asn Ser Thr Trp  
 355 360  
 25 AAT GTT ACT AAA GGG TTG AAT AAC ACT GAA GGA AAT ACC 1128  
 Asn Val Thr Lys Gly Leu Asn Asn Thr Glu Gly Asn Ser  
 365 370 375  
 30 ACA GGG GAT GAA AAT ATC ATA CTC CCA TGT AGA ATA AAA 1167  
 Thr Gly Asp Glu Asn Ile Ile Leu Pro Cys Arg Ile Lys  
 380 385  
 35 CAA ATT ATA AAC ATG TGG CAG GAA GCA GGA AAA GCA ATG 1206  
 Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys Ala Met  
 390 395 400  
 40 TAT GCC CCT CCC ATC AGT GGA CAA ATT AGA TGT TCA TCA 1245  
 Tyr Ala Pro Pro Ile Ser Gly Gln Ile Arg Cys Ser Ser  
 405 410 415  
 45 AAT ATT ACA GGG CTG CTA CTA ACA AGA GAT GGT GGT AGT 1284  
 Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly Ser  
 420 425  
 50 AAG AAC GAG AGC ATC ACC ACC GAG GTC TTC AGA CCT GGA 1323  
 Lys Asn Glu Ser Ile Thr Thr Glu Val Phe Arg Pro Gly  
 430 435 440  
 55 GGA GGA GAT ATG AGG GAC AAT TCG ACA AGT GAA TTA TAT 1362  
 Gly Gly Asp Met Arg Asp Asn Trp Arg Ser Glu Leu Tyr  
 445 450  
 60 AAA TAT AAA GTA GTA AAA ATT GAA CCA TTA GGA GCA GCG 1401  
 Lys Tyr Lys Val Val Lys Ile Glu Pro Leu Gly Val Ala  
 455 460 465  
 65 CCC ACC AAG GCA AAG AGA AGA GTG GTG CAG AGA GAA AAA 1440  
 Pro Thr Lys Ala Lys Arg Arg Val Val Gln Arg Glu Lys  
 470 475 480  
 70 AGA GCA GTG GGA ACA ATA GGA CCT ATG TTC CTT GGG TTC 1479  
 Arg Ala Val Gly Thr Ile Gly Ala Met Phe Leu Gly Phe  
 485 490  
 75 TTA GGA GCA TAA AGC TTC TAG A 1501  
 Leu Gly Ala Xaa Ser Phe Xaa  
 495 500

## (2) INFORMATION FOR SEQ ID NO:12:

## (1) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 500 amino acids

(B) TYPE: Amino Acid

5 (C) TOPOLOGY: Linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:12:

Leu	Glu	Val	Pro	Val	Trp	Lys	Glu	Ala	Thr	Thr	Thr	Leu	Phe	Cys	
1									10					15	
10	Ala	Ser	Asp	Ala	Lys	Ala	Tyr	Asn	Thr	Glu	Lys	His	Asn	Val	Trp
					20					25					30
15	Ala	Thr	His	Ala	Cys	Val	Pro	Thr	Asp	Pro	Asn	Pro	Gln	Glu	Val
					35					40					45
20	Val	Leu	Gly	Asn	Val	Thr	Glu	Asn	Phe	Asn	Met	Trp	Lys	Asn	Asn
					50					55					60
25	Met	Val	Glu	Gln	Met	His	Glu	Asp	Ile	Ile	Ser	Leu	Trp	Asp	Gln
					65					70					75
30	Ser	Leu	Lys	Pro	Cys	Val	Lys	Leu	Thr	Pro	Leu	Cys	Val	Thr	Leu
					80					85					90
35	Asn	Cys	Thr	Asp	Asp	Leu	Gly	Asn	Ala	Thr	Asn	Thr	Asn	Ser	Ser
					95					100					105
40	Ala	Thr	Thr	Asn	Ser	Ser	Ser	Trp	Glu	Glu	Met	Lys	Gly	Glu	Met
					110					115					120
45	Lys	Arg	Cys	Ser	Phe	Asn	Ile	Thr	Thr	Ser	Ile	Arg	Asp	Lys	Ile
					125					130					135
50	Lys	Lys	Glu	His	Ala	Leu	Phe	Tyr	Arg	Leu	Asp	Val	Val	Pro	Ile
					140					145					150
55	Asp	Asn	Asp	Asn	Thr	Thr	Tyr	Arg	Leu	Ile	Asn	Cys	Asn	Thr	Ser
					155					160					165
60	Val	Ile	Thr	Gln	Ala	Cys	Pro	Lys	Val	Ser	Phe	Glu	Pro	Ile	Pro
					170					175					180
65	Ile	His	Phe	Cys	Ala	Pro	Ala	Gly	Phe	Ala	Ile	Leu	Lys	Cys	Asn
					185					190					195
70	Asn	Lys	Thr	Phe	Glu	Gly	Lys	Gly	Pro	Cys	Lys	Asn	Val	Ser	Thr
					200					205					210
75	Val	Gln	Cys	Thr	His	Gly	Ile	Arg	Pro	Val	Val	Ser	Thr	Gln	Leu
					215					220					225
80	Leu	Leu	Asn	Gly	Ser	Leu	Ala	Glu	Glu	Val	Ile	Ile	Arg	Ser	
					230					235					240
85	Gly	Asn	Ile	Thr	Asp	Asn	Thr	Lys	Thr	Ile	Ile	Val	Gln	Leu	Asn
					245					250					255
90	Glu	Ser	Val	Val	Ile	Asn	Cys	Thr	Arg	Ser	Asn	Asn	Asn	Thr	Arg
					260					265					270
95	Lys	Ser	Ile	His	Ile	Gly	Pro	Gly	Ser	Ala	Phe	Phe	Ala	Thr	Gly
					275					280					285

Glu Ile Ile Gly Asp Ile Arg Gln Ala His Cys Asn Leu Ser Arg  
 290 295 300  
 5 Thr Gln Trp Asn Asn Thr Leu Gly Lys Ile Val Ile Lys Leu Arg  
 305 310 315  
 Glu Gln Phe Arg Lys Gln Phe Gly Glu Lys Thr Ile Val Phe Asn  
 320 325 330  
 10 Arg Ser Ser Gly Gly Asp Pro Glu Ile Ala Met His Ser Phe Asn  
 335 340 345  
 Cys Gly Gly Glu Phe Phe Tyr Cys Asn Thr Thr Ala Leu Phe Asn  
 350 355 360  
 15 Ser Thr Trp Asn Val Thr Lys Gly Leu Asn Asn Thr Glu Gly Asn  
 365 370 375  
 Ser Thr Gly Asp Glu Asn Ile Ile Leu Pro Cys Arg Ile Lys Gln  
 20 380 385 390  
 Ile Ile Asn Met Trp Gln Glu Val Gly Lys Ala Met Tyr Ala Pro  
 395 400 405  
 25 Pro Ile Ser Gly Gin Ile Arg Cys Ser Ser Asn Ile Thr Gly Leu  
 410 415 420  
 Leu Leu Thr Arg Asp Gly Gly Ser Lys Asn Glu Ser Ile Thr Thr  
 425 430 435  
 30 Glu Val Phe Arg Pro Gly Gly Asp Met Arg Asp Asn Trp Arg  
 440 445 450  
 Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile Glu Pro Leu Gly  
 35 455 460 465  
 Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Cln Arg Glu Lys  
 470 475 480  
 40 Arg Ala Val Gly Thr Ile Gly Ala Met Phe Leu Gly Phe Leu Gly  
 485 490 495  
 Ala Xaa Ser Phe Xaa  
 500  
 45 (2) INFORMATION FOR SEQ ID NO:13:  
 (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 1514 base pairs  
 (B) TYPE: Nucleic Acid  
 50 (C) STRANDEDNESS: Single  
 (D) TOPOLOGY: Linear  
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:13:  
 55 GG GAA TTC GGA TCC GGG GTA CCT GTG TGG AAG GAA GCA 38  
 Glu Phe Gly Ser Gly Val Pro Val Trp Lys Glu Ala  
 1 5 10  
 60 ACC ACC ACT CTA TTC TGT GCA TCA GAT CCT AGA GCA TAT 77  
 Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Arg Ala Tyr  
 15 20 25  
 GAC ACA GAG GTA CAT AAT GTT TGG GCC ACA CAT GCC TGT 116  
 Asp Thr Glu Val His Asn Val Trp Ala Thr His Ala Cys  
 65 30 35



1 AAT ACT AAA ACC ATA ATA GTA CAG CTG AAA GAA TCT GTA 779  
 Asn Thr Lys Thr Ile Ile Val Gln Leu Lys Glu Ser Val  
 250 255

5 GAA ATT AAT TGT ATA AGA CCC AAC AAT AAT ACA AGA AAA 818  
 Glu Ile Asn Cys Ile Arg Pro Asn Asn Asn Thr Arg Lys  
 260 265 270

10 GGT ATA CAT ATA GGA CCA GGG AGA GCA TGG TAT GCA ACA 857  
 Gly Ile His Ile Gly Pro Gly Ala Trp Tyr Ala Thr  
 275 280 285

15 GGA GAA ATA GTA GGA GAT ATA AGA AAG GCA TAT TGT AAC 896  
 Gly Glu Ile Val Gly Asp Ile Arg Lys Ala Tyr Cys Asn  
 290 295

20 ATT AGT AGA ACA AAA TGG AAT AAC ACT TTA ATA CAG ATA 935  
 Ile Ser Arg Thr Lys Trp Asn Asn Thr Leu Ile Gln Ile  
 300 305 310

25 GCT AAC AAA TTA AAA GAA AAA TAT AAT ACA ACA ATA AGC 974  
 Ala Asn Lys Leu Lys Glu Lys Tyr Asn Thr Thr Ile Ser  
 315 320

30 TTT AAT CGA TCC TCA GGA GGG GAC CCA GAA ATT GTA ACG 1013  
 Phe Asn Arg Ser Ser Gly Gly Asp Pro Glu Ile Val Thr  
 325 330 335

35 CAT AGT TTT AAT TGT GGA GGG GAG TTT TTC TAC TGT GAT 1052  
 His Ser Phe Asn Cys Gly Gly Phe Phe Tyr Cys Asp  
 340 345 350

40 TCA ACA CAA CTG TTT AAT AGT ACT TGG AAT TTA AAT GGT 1091  
 Ser Thr Gln Leu Phe Asn Ser Thr Trp Asn Leu Asn Gly  
 355 360

45 ACT TGG AAT TTT ACT GCA GGG TCA AAT GAA ACT GAA GGC 1130  
 Thr Trp Asn Phe Thr Ala Gly Ser Asn Glu Thr Glu Gly  
 365 370 375

50 AAT ATC ACA CTC CCA TGC AGA ATA AAA CAA ATT ATA AAC 1169  
 Asn Ile Thr Leu Pro Cys Arg Ile Lys Gln Ile Ile Asn  
 380 385

55 AGG TGG CAG GAA GTA GGG AAA GCA ATG TAT GCC CCT CCC 1208  
 Arg Trp Gln Glu Val Gly Lys Ala Met Tyr Ala Pro Pro  
 390 395 400

60 ATC AGT GGA CAA ATA AAA TGC TCA TCA AAC ATT ACA GGG 1247  
 Ile Ser Gly Gln Ile Lys Cys Ser Ser Asn Ile Thr Gly  
 405 410 415

65 GAG AGC AGT ACT ACT GAG ACC TTC AGA CCG GGA GGA GGA 1325  
 Glu Ser Ser Thr Thr Glu Thr Phe Arg Pro Gly Gly Gly  
 430 435 440

70 GAT ATG AGG AAC AAT TGG AGA AGT GAA TTA TAT AAA TAT 1364  
 Asp Met Arg Asn Asn Trp Arg Ser Glu Leu Tyr Lys Tyr  
 445 450

AAA GTC GTC AAA ATT GAA CCA TTA GGA GTC GCA CCC ACC 1403  
 Lys Val Val Lys Ile Glu Pro Leu Gly Val Ala Pro Thr  
 455 460 465  
 5 AAG GCA AAG AGA AGA GTG GTG CAG AGA GAA AAA AGA GCA 1442  
 Lys Ala Lys Arg Arg Val Val Gln Arg Glu Lys Arg Ala  
 470 475 480  
 GTG GGA GCG CTA GGA GCT ATG TTC CTT GGG TTC TTA GGA 1481  
 Val Gly Ala Leu Gly Ala Met Phe Leu Gly Phe Leu Gly  
 485 490  
 GCA TAA AGC TTC TAG ACC GAC TCT AGA GGA TCC 1514  
 Ala Xaa Ser Phe Xaa Thr Asp Ser Arg Gly Ser  
 15 495 500 504

## (2) INFORMATION FOR SEQ ID NO:14:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 504 amino acids
- (B) TYPE: Amino Acid
- (C) TOPOLOGY: Linear
- (xi) SEQUENCE DESCRIPTION: SEQ. ID NO:14:

Glu Phe Gly Ser Gly Val Pro Val Trp Lys Glu Ala Thr Thr Thr  
 25 1 5 10 15  
 Leu Phe Cys Ala Ser Asp Ala Arg Ala Tyr Asp Thr Glu Val His  
 20 20 25 30  
 Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Ser Pro  
 30 35 40 45  
 Gln Glu Val Val Leu Glu Asn Val Thr Glu Asn Phe Asn Met Trp  
 35 50 55 60  
 Lys Asn Asn Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu  
 40 65 70 75  
 Trp Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys  
 45 80 85 90  
 Val Thr Leu Asn Cys Ser Asp Tyr Arg Asn Ala Thr Asp Tyr Lys  
 50 95 100 105  
 Asn Ala Thr Asp Thr Thr Ser Ser Asn Glu Gly Lys Met Glu Arg  
 55 110 115 120  
 Gly Glu Ile Lys Asn Cys Ser Phe Asn Ile Thr Thr Ser Ile Lys  
 60 125 130 135  
 Asn Lys Met Gln Lys Glu Tyr Ala Leu Phe Tyr Lys Leu Asp Ile  
 65 140 145 150  
 Val Pro Ile Asp Asn Thr Ser Tyr Thr Leu Ile Ser Cys Asn Thr  
 70 155 160 165  
 Ser Val Ile Thr Gln Ala Cys Pro Lys Val Ser Phe Glu Pro Thr  
 75 170 175 180  
 Pro Ile His Tyr Cys Ala Pro Ala Gly Phe Ala Ile Leu Lys Cys  
 80 185 190 195  
 Asn Asp Lys Lys Phe Ser Gly Lys Glu Cys Lys Asn Val Ser  
 85 200 205 210  
 90

	Thr	Val	Gln	Cys	Thr	His	Gly	Ile	Arg	Pro	Val	Val	Ser	Thr	Gln
	215													220	225
5	Leu	Leu	Leu	Asn	Gly	Ser	Leu	Ala	Glu	Glu	Glu	Val	Val	Ile	Arg
					230				235					240	
	Ser	Asp	Asn	Phe	Ile	Asp	Asn	Thr	Lys	Thr	Ile	Ile	Val	Gln	Leu
					245				250					255	
10	Lys	Glu	Ser	Val	Glu	Ile	Asn	Cys	Ile	Arg	Pro	Asn	Asn	Asn	Thr
					260				265					270	
	Arg	Lys	Gly	Ile	His	Ile	Gly	Pro	Gly	Arg	Ala	Trp	Tyr	Ala	Thr
					275				280					285	
15	Gly	Glu	Ile	Val	Gly	Asp	Ile	Arg	Lys	Ala	Tyr	Cys	Asn	Ile	Ser
					290				295					300	
20	Arg	Thr	Lys	Trp	Asn	Asn	Thr	Leu	Ile	Gln	Ile	Ala	Asn	Lys	Leu
					305				310					315	
	Lys	Glu	Lys	Tyr	Asn	Thr	Thr	Ile	Ser	Phe	Asn	Arg	Ser	Ser	Gly
					320				325					330	
25	Gly	Asp	Pro	Glu	Ile	Val	Thr	His	Ser	Phe	Asn	Cys	Gly	Gly	Glu
					335				340					345	
	Phe	Phe	Tyr	Cys	Asp	Ser	Thr	Gln	Leu	Phe	Asn	Ser	Thr	Trp	Asn
					350				355					360	
30	Leu	Asn	Gly	Thr	Trp	Asn	Phe	Thr	Ala	Gly	Ser	Asn	Glu	Thr	Glu
					365				370					375	
	Gly	Asn	Ile	Thr	Leu	Pro	Cys	Arg	Ile	Lys	Gln	Ile	Ile	Asn	Arg
					380				385					390	
	Trp	Gln	Glu	Val	Gly	Lys	Ala	Met	Tyr	Ala	Pro	Pro	Ile	Ser	Gly
					395				400					405	
40	Gln	Ile	Lys	Cys	Ser	Ser	Asn	Ile	Thr	Gly	Met	Ile	Leu	Thr	Arg
					410				415					420	
	Asp	Gly	Gly	Asn	Glu	Asn	Asn	Glu	Ser	Ser	Thr	Thr	Glu	Thr	
					425				430					435	
45	Phe	Arg	Pro	Gly	Gly	Asp	Met	Arg	Asn	Asn	Trp	Arg	Ser	Glu	
					440				445					450	
	Leu	Tyr	Lys	Tyr	Lys	Val	Val	Lys	Ile	Glu	Pro	Leu	Gly	Val	Ala
					455				460					465	
	Pro	Thr	Lys	Ala	Lys	Arg	Arg	Val	Val	Gln	Arg	Gl	Lys	Arg	Ala
					470				475					480	
55	Val	Gly	Ala	Leu	Gly	Ala	Met	Phe	Leu	Gly	Phe	Leu	Gly	Ala	Xaa
					485				490					495	
	Ser	Phe	Xaa	Thr	Asp	Ser	Arg	Gly	Ser						
					500				504						

(2) INFORMATION FOR SEQ ID NO:15:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1408 base pairs

(B) TYPE: Nucleic Acid

(C) STRANDEDNESS: Single

## (D) TOPOLOGY: Linear

## (xi) SEQUENCE DESCRIPTION: SEQ\_ID NO:15:

5	G	CTA	CCT	GTG	TGG	AAG	GAA	GCA	ACC	ACC	ACT	CTA	TTC	37
	Val	Pro	Val	Trp	Lys	Glu	Ala	Thr	Thr	Thr	Leu	Phe		
	1				5					10				
10	TGT	GCA	TCA	GAT	GCT	AGA	GCA	TAT	GAC	ACA	GAG	GTA	CAT	76
	Cys	Ala	Ser	Asp	Ala	Arg	Ala	Tyr	Asp	Thr	Glu	Val	His	
	15				20						25			
15	AAT	GTT	TGG	GCC	ACA	CAT	GCC	TGT	GTA	CCC	ACA	GAC	CCT	115
	Asn	Val	Trp	Ala	Thr	His	Ala	Cys	Val	Pro	Thr	Asp	Pro	
	30				35									
20	AGT	CCA	CAA	GAA	GTA	TTT	TTG	GGA	AAT	GTG	ACA	GAA	AAT	154
	Ser	Pro	Gln	Glu	Val	Phe	Leu	Gly	Asn	Val	Thr	Glu	Asn	
	40				45						50			
25	TTT	AAT	ATG	TGG	AAA	AAT	AAC	ATG	GTA	GAA	CAA	ATG	TAT	193
	Phe	Asn	Met	Trp	Lys	Asn	Asn	Met	Val	Glu	Gln	Met	Tyr	
	55				60									
30	GAG	GAT	ATA	ATT	AGT	TTA	TGG	GAT	CAA	AGC	TTA	AAG	CCA	232
	Glu	Asp	Ile	Ile	Ser	Leu	Trp	Asp	Gln	Ser	Leu	Lys	Pro	
	65				70						75			
35	TGT	GTA	AAA	TTA	ACC	CCA	CTC	TGT	GTT	ACT	TTA	AAT	TGC	271
	Cys	Val	Lys	Leu	Thr	Pro	Leu	Cys	Val	Thr	Leu	Asn	Cys	
	80				85						90			
40	AGT	GAT	TAT	AGG	AAT	GCT	ACT	GAT	TAT	AAG	AAT	CCT	ACT	310
	Ser	Asp	Tyr	Arg	Asn	Ala	Thr	Asp	Tyr	Lys	Asn	Ala	Thr	
	95				100									
45	GAT	ACC	ACT	AGT	AAC	GAG	GGA	AAG	ATG	GAG	AGA	CCA	349	
	Asp	Thr	Ser	Ser	Asn	Glu	Gly	Lys	Met	Glu	Arg	Gly		
	105				110					115				
50	GAA	ATA	AAA	AAC	TGC	TCT	TTC	AAT	ATC	ACC	ACA	AGC	ATA	388
	Glu	Ile	Lys	Cys	Ser	Phe	Asn	Ile	Thr	Thr	Ser	Ile		
	120				125									
55	AAA	AAT	AAG	ATG	CAG	AAA	GAA	TAT	GCA	CTT	TTC	TAT	AAA	427
	Lys	Asn	Lys	Met	Gln	Lys	Glu	Tyr	Ala	Leu	Phe	Tyr	Lys	
	130				135					140				
60	CTT	AAT	ATA	GTA	CCA	ATA	GAT	AAT	ACA	AGC	TAT	ACA	TTG	466
	Leu	Asn	Ile	Val	Pro	Ile	Asp	Asn	Thr	Ser	Tyr	Thr	Leu	
	145				150					155				
65	ATA	AGT	TGT	AAC	ACC	TCA	GTC	ATT	ACA	CAG	GCC	TGT	CCA	505
	Ile	Ser	Cys	Asn	Thr	Ser	Val	Ile	Thr	Gln	Ala	Cys	Pro	
	160				165									
70	AAG	GTA	TCC	TTT	GAA	CCA	ATT	CCC	ATA	CAT	TAT	TGT	GCT	544
	Lys	Val	Ser	Phe	Glu	Pro	Ile	Pro	Ile	His	Tyr	Cys	Ala	
	170				175					180				
75	CCG	GCT	GGT	TTT	GCG	ATT	CTA	AAG	TGT	AAT	GAT	AAG	AAG	583
	Pro	Ala	Gly	Phe	Ala	Ile	Leu	Lys	Cys	Asn	Asp	Lys	Lys	
	185				190									

195            TTC AGT GGA AAA GGA GAA TGT AAA AAT CTC AGC ACA GTC 622  
 Phe Ser Gly Lys Gly Glu Cys Lys Asn Val Ser Thr Val  
 200            205  
 5            CAA TGT ACA CAT GGA ATT AGG CCA GTC GTC TCA ACT CAA 661  
 Gln Cys Thr His Gly Ile Arg Pro Val Val Ser Thr Gln  
 210            215            220  
 10            CTG CTG TTA AAT GGC AGT CTC GCA GAA GAA GAG GTG GTA 700  
 Leu Leu Leu Asn Gly Ser Leu Ala Glu Glu Glu Val Val  
 225            230  
 15            ATT AGA TCT GAC AAT TTC ACA GAC AAT ACT AAA ACC ATA 739  
 Ile Arg Ser Asp Asn Phe Thr Asp Asn Thr Lys Thr Ile  
 235            240            245  
 20            ATA GTA CAG CTG AAA GAA TCT GTC GAA ATT AAT TGT ATA 778  
 Ile Val Gln Leu Lys Glu Ser Val Glu Ile Asn Cys Ile  
 250            255  
 25            AGA CCC AAC AAT AAT ACA AGA AAA GGT ATA CAT ATA GGA 817  
 Arg Pro Asn Asn Asn Thr Arg Lys Gly Ile His Ile Gly  
 260            265            270  
 30            CCA GGG AGA GCA TGG TAT GCA ACA GGA GAA ATA GTA GGA 856  
 Pro Gly Arg Ala Trp Tyr Ala Thr Gly Glu Ile Val Gly  
 275            280            285  
 35            GAT ATA AGA CAG GCA TAT TGT AAC ATT AGT AGA ACA AAA 895  
 Asp Ile Arg Gln Ala Tyr Cys Asn Ile Ser Arg Thr Lys  
 290            295  
 40            TGG AAT AAC ACT TTA ATA CAG ATA GCT AAC AAA TTA AAA 934  
 Trp Asn Asn Thr Leu Ile Gln Ile Ala Asn Lys Leu Lys  
 300            305            310  
 45            GAA AAA TAT AAT ACA ACA ATA ACC TTT AAT CGA TCC TCA 973  
 Glu Lys Tyr Asn Thr Thr Ile Ser Phe Asn Arg Ser Ser  
 315            320  
 50            GGA GGG GAC CCA GAA ATT GTC ACC CAT ACT TTT AAT TGT 1012  
 Gly Gly Asp Pro Glu Ile Val Thr His Ser Phe Asn Cys  
 325            330            335  
 55            GGA GGG GAA TTT TTC TAC TGT AAT TCA ACA CAA CTG TTT 1051  
 Gly Gly Glu Phe Phe Tyr Cys Asn Ser Thr Gln Leu Phe  
 340            345            350  
 60            AAT AGT ACT TGG AAT TTA AAT GGT ACT TGG AAT TTT ACT 1090  
 Asn Ser Thr Trp Asn Leu Asn Gly Thr Trp Asn Phe Thr  
 355            360  
 65            GCA GGG TCA AAT GAA ACT GAA GGC AAT ATC ACA CTC CCA 1129  
 Ala Gly Ser Asn Glu Thr Glu Gly Asn Ile Thr Leu Pro  
 365            370            375  
 70            TGC AGA ATA AAA CAA ATT ATA AAC ACC TGG CAG GAA GTA 1168  
 Cys Arg Ile Lys Gln Ile Ile Asn Arg Trp Gln Glu Val  
 380            385  
 75            GGA AAA GCA ATG TAT GCC CCT CCC ATC AGT GGA CAA ATA 1207  
 Gly Lys Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile  
 390            395            400

AGA TGC TCA TCA AAC ATT ACA GGG ATG ATA TTA ACA AGG 1246  
 Arg Cys Ser Ser Asn Ile Thr Gly Met Ile Leu Thr Arg  
 405 410 415  
 5. GAT GGT GGT AAC GAG AAC AAT AAT GAG AGC AGT ACT ACT 1285  
 Asp Gly Gly Asn Glu Asn Asn Asn Glu Ser Ser Thr Thr  
 420 425  
 10. GAG ACC TTC AGA CCG GGA GGA GGA GAT ATG AGG AAC AAT 1324  
 Glu Thr Phe Arg Pro Gly Gly Asp Met Arg Asn Asn  
 430 435 440  
 15. TGG AGA AGT GAA TTA TAT AAA TAT AAA GTA GTA AAA ATT 1363  
 Trp Arg Ser Glu Leu Tyr Lys Tyr Val Val Lys Ile  
 445 450  
 20. GAG CCA TTA GGA GTA GCA CCC ACC GAC TCT AGA GGA TCC 1402  
 Glu Pro Leu Gly Val Ala Pro Thr Asp Ser Arg Gly Ser  
 455 460 465  
 25. TCT AGA 1408  
 Ser Arg  
 469

25. (2) INFORMATION FOR SEQ ID NO:16:  
 (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 469 amino acids  
 (B) TYPE: Amino Acid  
 (D) TOPOLOGY: Linear  
 30. (xi) SEQUENCE DESCRIPTION: SEQ ID NO:16:  
 Val Pro Val Trp Lys Glu Ala Thr Thr Leu Phe Cys Ala Ser  
 1 5 10 15  
 35. Asp Ala Arg Ala Tyr Asp Thr Glu Val His Asn Val Trp Ala Thr  
 20 25 30  
 His Ala Cys Val Pro Thr Asp Pro Ser Pro Gln Glu Val Phe Leu  
 35 40 45  
 40. Gly Asn Val Thr Glu Asn Phe Asn Met Trp Lys Asn Asn Met Val  
 50 55 60  
 Glu Gln Met Tyr Glu Asp Ile Ile Ser Leu Trp Asp Gln Ser Leu  
 65 70 75  
 45. Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu Asn Cys  
 80 85 90  
 50. Ser Asp Tyr Arg Asn Ala Thr Asp Tyr Lys Asn Ala Thr Asp Thr  
 95 100 105  
 Thr Ser Ser Asn Glu Gly Lys Met Glu Arg Gly Glu Ile Lys Asn  
 110 115 120  
 55. Cys Ser Phe Asn Ile Thr Thr Ser Ile Lys Asn Lys Met Gln Lys  
 125 130 135  
 60. Glu Tyr Ala Leu Phe Tyr Lys Leu Asn Ile Val Pro Ile Asp Asn  
 140 145 150  
 Thr Ser Tyr Thr Leu Ile Ser Cys Asn Thr Ser Val Ile Thr Gln  
 155 160 165

	Ala	Cys	Pro	Lys	Val	Ser	Phe	Glu	Pro	Ile	Pro	Ile	His	Tyr	Cys
					170										180
5	Ala	Pro	Ala	Gly	Phe	Ala	Ile	Leu	Lys	Cys	Asn	Asp	Lys	Lys	Phe
					185					190					195
10	Ser	Gly	Lys	Gly	Glu	Cys	Lys	Asn	Val	Ser	Thr	Val	Gln	Cys	Thr
					200					205					210
15	His	Gly	Ile	Arg	Pro	Val	Val	Ser	Thr	Gln	Leu	Leu	Leu	Asn	Gly
					215					220					225
20	Ser	Leu	Ala	Glu	Glu	Glu	Val	Val	Ile	Arg	Ser	Asp	Asn	Phe	Thr
					230					235					240
25	Asp	Asn	Thr	Lys	Thr	Ile	Ile	Val	Gln	Leu	Lys	Glu	Ser	Val	Glu
					245					250					255
30	Ile	Asn	Cys	Ile	Arg	Pro	Asn	Asn	Asn	Thr	Arg	Lys	Gly	Ile	His
					260					265					270
35	Ile	Gly	Pro	Gly	Arg	Ala	Trp	Tyr	Ala	Thr	Gly	Glu	Ile	Vai	Gly
					275					280					285
40	Asp	Ile	Arg	Gln	Ala	Tyr	Cys	Asn	Ile	Ser	Arg	Thr	Lys	Trp	Asn
					290					295					300
45	Asn	Thr	Leu	Ile	Gln	Ile	Ala	Asn	Lys	Leu	Lys	Glu	Lys	Tyr	Asn
					305					310					315
50	Thr	Thr	Ile	Ser	Phe	Asn	Arg	Ser	Ser	Gly	Gly	Asp	Pro	Glu	Ile
					320					325					330
55	Val	Thr	His	Ser	Phe	Asn	Cys	Gly	Gly	Glu	Phe	Phe	Tyr	Cys	Asn
					335					340					345
60	Ser	Thr	Gln	Leu	Phe	Asn	Ser	Thr	Trp	Asn	Leu	Asn	Gly	Thr	Trp
					350					355					360
65	Asn	Phe	Thr	Ala	Gly	Ser	Asn	Glu	Thr	Glu	Gly	Asn	Ile	Thr	Leu
					365					370					375
70	Pro	Cys	Arg	Ile	Lys	Gln	Ile	Ile	Asn	Arg	Trp	Gln	Glu	Val	Gly
					380					385					390
75	Lys	Ala	Met	Tyr	Ala	Pro	Pro	Ile	Ser	Gly	Gln	Ile	Arg	Cys	Ser
					395					400					405
80	Ser	Asn	Ile	Thr	Gly	Met	Ile	Leu	Thr	Arg	Asp	Gly	Gly	Asn	Glu
					410					415					420
85	Asn	Asn	Asn	Glu	Ser	Ser	Thr	Thr	Glu	Thr	Phe	Arg	Pro	Gly	Gly
					425					430					435
90	Gly	Asp	Met	Arg	Asn	Asn	Trp	Arg	Ser	Glu	Leu	Tyr	Lys	Tyr	Lys
					440					445					450
95	Val	Val	Lys	Ile	Glu	Pro	Leu	Gly	Val	Ala	Pro	Thr	Asp	Ser	Arg
					455					460					465
100	Gly	Ser	Ser	Arg											
					469										

(2) INFORMATION FOR SEQ ID NO:17:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1499 base pairs

(B) TYPE: Nucleic Acid

(C) STRANDEDNESS: Single

(D) TOPOLOGY: Linear

5 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:17:

GAG GTA CCT GTG TGG AAA GAA GCA ACC ACT ACT CTA 36  
 Glu Val Pro Val Trp Lys Glu Ala Thr Thr Thr Leu  
 1 5 10

10 TTT TGT GCA TCA GAT GCT AAA GCA TAT GAC ACA GGG GTG 75  
 Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Gly Val  
 15 20 25

15 CAT AAT GTT TGG GCC ACA CAT GCC TGT GTA CCC ACA GAC 114  
 His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp  
 30 35

20 CCC AAC CCA CAA GAA ATA GAA TTG GTA AAT GTG ACA GAA 153  
 Pro Asn Pro Gln Glu Ile Glu Leu Val Asn Val Thr Glu  
 40 45 50

25 GAT TTT AAC ATG TGG AAA AAT AAA ATG GTA GAC CAG ATG 192  
 Asp Phe Asn Met Trp Lys Asn Lys Met Val Asp Gln Met  
 55 60

30 CAT GAG GAT ATA ATC AGT TTA TGG GAT GAA AGC CTA AAG 231  
 His Glu Asp Ile Ile Ser Leu Trp Asp Glu Ser Leu Lys  
 65 70 75

35 CCA TGT GTA AAC TTA ACC CCA CTT TGT GTT ACT CTA AAC 270  
 Pro Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu Asn  
 80 85 90

40 TGC AGT GAT GTG AAC AAT TCC ACA AAT CCT AAT GAT ACT 309  
 Cys Ser Asp Val Asn Asn Ser Thr Asn Pro Asn Asp Thr  
 95 100

45 AAT ACT AAT TCC ACT AAT ACT ACT TCC TCT ACT CCT ACG 348  
 Asn Thr Asn Ser Thr Asn Thr Ser Ser Thr Pro Thr  
 105 110 115

50 GCC ACT ACT AGT AGC GAG GAA AAG ATG GAC AAG GGA GAA 387  
 Ala Thr Thr Ser Ser Glu Glu Lys Met Glu Lys Gly Glu  
 120 125

55 ATA AAA AAC TGC TCT TTC AAT ATC ACC ACA CAC ATG AAA 426  
 Ile Lys Asn Cys Ser Phe Asn Ile Thr Thr His Met Lys  
 130 135 140

60 GAT AAG GCA CAG AAA GAA TAT GCA CTT TTT TAT AAA CTT 465  
 Asp Lys Ala Gln Lys Glu Tyr Ala Leu Phe Tyr Lys Leu  
 145 150 155

65 GAT ATA GTA CCA ATA GAT GAT AAT AAT GCC AGC TAT AGG 504  
 Asp Ile Val Pro Ile Asp Asp Asn Asn Ala Ser Tyr Arg  
 160 165

70 TTG ATA AGT TGT AAT ACC TCA GAC ATT ACA CAG GCC TGT 543  
 Leu Ile Ser Cys Asn Thr Ser Asp Ile Thr Gln Ala Cys  
 170 175 180

75 CCA AAG GTG ACC TTT GAG CCA ATT CCC ATA CAT TAT TGT 582  
 Pro Lys Val Thr Phe Glu Pro Ile Pro Ile His Tyr Cys  
 185 190

195      GCC CCG GCT GGT TTT GCG ATT CTA AAC TGT AAA GAT AAG 621  
 Ala Pro Ala Gly Phe Ala Ile Leu Lys Cys Lys Asp Lys  
 200      205  
 5      AAG TTC AAT CGA ACA GGA CCA TGT TCA AAG GTC AGC ACA 660  
 Lys Phe Asn Gly Thr Gly Pro Cys Ser Lys Val Ser Thr  
 210      215      220  
 10      GTA CAA TGT ACA CAT GGA ATT AGG CCA GTA GTA TCA ACT 699  
 Val Gln Cys Thr His Gly Ile Arg Pro Val Val Ser Thr  
 225      230  
 15      CAA CTG TTG TTA AAT GGC AGT CTT GCA GAA GAA GAA GTA 738  
 Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu Glu Glu Val  
 235      240      245  
 20      GTA ATT AGA TCT GTC AAT TTC ACA GAC AAT GCT AAA ATC 777  
 Val Ile Arg Ser Val Asn Phe Thr Asp Asn Ala Lys Ile  
 250      255  
 25      ATA ATA GTA CAG CTG AAA GAA CCT GTA GCA ATT AAT TGT 816  
 Ile Ile Val Gln Leu Lys Glu Pro Val Ala Ile Asn Cys  
 260      265      270  
 30      ACA AGA CCC AAC AAC AAT ACA AGA AAA GGT ATA CAT CTA 855  
 Thr Arg Pro Asn Asn Asn Thr Arg Lys Gly Ile His Leu  
 275      280      285  
 35      GGA CCA GGG AGC ACA TTT TAT ACA ACA GGA GAA ATA ATA 894  
 Gly Pro Gly Ser Thr Phe Tyr Thr Thr Gly Glu Ile Ile  
 290      295  
 40      GGA GAC ATA AGA AAA GCA TAT TGC AAG ATT AGT AAA GAA 933  
 Gly Asp Ile Arg Lys Ala Tyr Cys Lys Ile Ser Lys Glu  
 300      305      310  
 45      AAA TGG AAT AAC ACT TTA AGA CAG GTA GTT AAA AAA TTA 972  
 Lys Trp Asn Asn Thr Leu Arg Gln Val Val Lys Lys Leu  
 315      320  
 50      AGA GAA CAA TTT GGG AAT AAA ACA ATA ATT TTT AAT CGA 1011  
 Arg Glu Gln Phe Gly Asn Lys Thr Ile Ile Phe Asn Arg  
 325      330      335  
 55      TCC TCA GGA GGG GAC CCA GAA ATT GTA ATG CAC AGT TTT 1050  
 Ser Ser Gly Gly Asp Pro Glu Ile Val Met His Ser Phe  
 340      345      350  
 60      AAC TGT GGA GGG GAG TTT TTC TAC TGT AAT ACA ACA CAA 1089  
 Asn Cys Gly Gly Glu Phe Phe Tyr Cys Asn Thr Thr Gln  
 355      360  
 65      CTG TTT AAT AGT ACT TGG AAT AAT ACT GAA GGG ACA AAT 1128  
 Leu Phe Asn Ser Thr Trp Asn Asn Thr Glu Gly Thr Asn  
 365      370      375  
 70      AGC ACT GAA GGA AAT AGC ACA ATC ACA CTC CCA TGC AGA 1167  
 Ser Thr Glu Gly Asn Ser Thr Ile Thr Leu Pro Cys Arg  
 380      385  
 75      ATA AAA CAA ATT ATA AAT ATG TGG CAG GAA GTA GGA AAA 1206  
 Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys  
 390      395      400

GCA ACG TAT GCC CCT CCC ATC AGA GGA CGA ATT AGA TGC 1245  
 Ala Thr Tyr Ala Pro Pro Ile Arg Gly Arg Ile Arg Cys  
 405 410 415  
 5 ATA TCA AAT ATT ACA GGA CTG CTA TTA ACA AGA GAT GGT 1284  
 Ile Ser Asn Ile Thr Gly Leu Leu Thr Arg Asp Gly  
 420 425  
 10 GGT AGG AAT GTC ACA AAC AAT ACC GAA ACC TTC AGA CCT 1323  
 Gly Arg Asn Val Thr Asn Asn Thr Glu Thr Phe Arg Pro  
 430 435 440  
 15 GGA CGA GGA GAC ATG AGG GAC AAT TGG AGA AGT GAA TTA 1362  
 Gly Gly Gly Asp Met Arg Asp Asn Trp Arg Ser Glu Leu  
 445 450  
 20 TAT AAA TAT AAA GTA GTA AAA CTT GAA CCA TTA GGA ATA 1401  
 Tyr Lys Tyr Lys Val Val Lys Val Glu Pro Leu Gly Ile  
 455 460 465  
 25 GCA CCC ACC AAG GCA AAG AGA AGA GTG GTG CAC AGA GAC 1440  
 Ala Pro Thr Lys Ala Lys Arg Arg Val Val His Arg Asp  
 470 475 480  
 30 AAA AGA GCA GCA CTA GGA GCC TTG TTC CTT GGG TTC TTA 1479  
 Lys Arg Ala Ala Leu Gly Ala Leu Phe Leu Gly Phe Leu  
 485 490  
 GGA GCA TAA AAG CTT CTA GA 1499  
 35 Gly Ala Xaa Lys Leu Leu  
 495 499  
 (2) INFORMATION FOR SEQ ID NO:18:  
 (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 499 amino acids  
 (B) TYPE: Amino Acid  
 (D) TOPOLOGY: Linear  
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:18:  
 40 Glu Val Pro Val Trp Lys Glu Ala Thr Thr Leu Phe Cys Ala  
 1 5 10 15  
 Ser Asp Ala Lys Ala Tyr Asp Thr Gly Val His Asn Val Trp Ala  
 20 25 30  
 45 Thr His Ala Cys Val Pro Thr Asp Pro Asn Pro Gln Glu Ile Glu  
 35 40 45  
 Leu Val Asn Val Thr Glu Asp Phe Asn Met Trp Lys Asn Lys Met  
 50 55 60  
 Val Asp Gln Met His Glu Asp Ile Ile Ser Leu Trp Asp Glu Ser  
 65 70 75  
 55 Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu Asn  
 80 85 90  
 Cys Ser Asp Val Asn Asn Ser Thr Asn Pro Asn Asp Thr Asn Thr  
 95 100 105  
 60 Asn Ser Thr Asn Thr Ser Ser Thr Pro Thr Ala Thr Thr Ser  
 110 115 120  
 Ser Glu Glu Lys Met Glu Lys Gly Glu Ile Lys Asn Cys Ser Phe  
 65 125 130 135

	Asn	Ile	Thr	Thr	His	Met	Lys	Asp	Lys	Ala	Gln	Lys	Glu	Tyr	Ala
						140				145					150
5	Leu	Phe	Tyr	Lys	Leu	Asp	Ile	Val	Pro	Ile	Asp	Asp	Asn	Asn	Ala
						155				160					165
10	Ser	Tyr	Arg	Leu	Ile	Ser	Cys	Asn	Thr	Ser	Asp	Ile	Thr	Gln	Ala
						170				175					180
15	Cys	Pro	Lys	Val	Thr	Phe	Glu	Pro	Ile	Pro	Ile	His	Tyr	Cys	Ala
						185				190					195
20	Pro	Ala	Gly	Phe	Ala	Ile	Leu	Lys	Cys	Lys	Asp	Lys	Lys	Phe	Asn
						200				205					210
25	Gly	Thr	Gly	Pro	Cys	Ser	Lys	Val	Ser	Thr	Val	Gln	Cys	Thr	His
						215				220					225
30	Gly	Ile	Arg	Pro	Val	Val	Ser	Thr	Gln	Leu	Leu	Leu	Asn	Gly	Ser
						230				235					240
35	Leu	Ala	Glu	Glu	Glu	Val	Val	Ile	Arg	Ser	Val	Asn	Phe	Thr	Asp
						245				250					255
40	Asn	Ala	Lys	Ile	Ile	Ile	Val	Gln	Leu	Lys	Glu	Pro	Val	Ala	Ile
						260				265					270
45	Asn	Cys	Thr	Arg	Pro	Asn	Asn	Asn	Thr	Arg	Lys	Gly	Ile	His	Leu
						275				280					285
50	Gly	Pro	Gly	Ser	Thr	Phe	Tyr	Thr	Gly	Glu	Ile	Ile	Gly	Asp	
						290				295					300
55	Ile	Arg	Lys	Ala	Tyr	Cys	Lys	Ile	Ser	Lys	Glu	Lys	Trp	Asn	Asn
						305				310					315
60	Thr	Leu	Arg	Gln	Val	Val	Lys	Lys	Leu	Arg	Glu	Gln	Phe	Gly	Asn
						320				325					330
65	Lys	Thr	Ile	Ile	Phe	Asn	Arg	Ser	Ser	Gly	Gly	Asp	Pro	Glu	Ile
						335				340					345
70	Val	Met	His	Ser	Phe	Asn	Cys	Gly	Gly	Glu	Phe	Phe	Tyr	Cys	Asn
						350				355					360
75	Thr	Thr	Gln	Leu	Phe	Asn	Ser	Thr	Trp	Asn	Asn	Thr	Glu	Gly	Thr
						365				370					375
80	Asn	Ser	Thr	Glu	Gly	Asn	Ser	Thr	Ile	Thr	Leu	Pro	Cys	Arg	Ile
						380				385					390
85	Lys	Gln	Ile	Ile	Asn	Met	Trp	Gln	Glu	Val	Gly	Lys	Ala	Thr	Tyr
						395				400					405
90	Ala	Pro	Pro	Ile	Arg	Gly	Arg	Ile	Arg	Cys	Ile	Ser	Asn	Ile	Thr
						410				415					420
95	Gly	Leu	Leu	Leu	Thr	Arg	Asp	Gly	Gly	Arg	Asn	Val	Thr	Asn	Asn
						425				430					435
100	Thr	Glu	Thr	Phe	Arg	Pro	Gly	Gly	Asp	Met	Arg	Asp	Asn	Trp	
						440				445					450
105	Arg	Ser	Glu	Leu	Tyr	Lys	Tyr	Lys	Val	Val	Lys	Val	Glu	Pro	Leu
						455				460					465

Gly Ile Ala Pro Thr Lys Ala Lys Arg Arg Val Val His Arg Asp  
470 475 480

5 Lys Arg Ala Ala Leu Gly Ala Leu Phe Leu Gly Phe Leu Gly Ala  
485 490 495

Xaa Lys Leu Leu  
499

10 (2) INFORMATION FOR SEQ ID NO:19:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1499 base pairs

(B) TYPE: Nucleic Acid

(C) STRANDEDNESS: Single

15 (D) TOPOLOGY: Linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:19:

GAG GTA CCT GTÀ TGG AAA GAA GCA ACC ACT ACT CTA 36  
Glu Val Pro Val Trp Lys Glu Ala Thr Thr Thr Leu  
20 1 5 10

TTT TGT GCA TCA GAT GCT AAA GCA TAT GAC ACA GAG GTG 75  
Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu Val  
25 15 20 25

25 CAT AAT GTT TGG GCC ACA CAT GCC TGT GTA CCC ACA GAC 114  
His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp  
30 35

30 CCC AAC CCA CAA GAA ATA GAA TTG GTA AAT GTG ACA GAA 153  
Pro Asn Pro Cln Glu Ile Glu Leu Val Asn Val Thr Glu  
40 45 50

35 GAT TTT AAC ATG TGG AAA AAT AAA ATG GTA GAC CAG ATG 192  
Asp Phe Asn Met Trp Lys Asn Lys Met Val Asp Cln Met  
55 60

40 CAT GAG GAT ATA ATC AGT TTA TGG GAT GAA AGC CTA AAG 231  
His Glu Asp Ile Ile Ser Leu Trp Asp Glu Ser Leu Lys  
65 70 75

45 CCA TGT GTA AAG TTA ACC CCA CTT TGT GTT ACT CTA AAC 270  
Pro Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu Asn  
80 85 90

50 TGC AGT GAT GTG AAC AAT TCC ACA AAT CCT AAT GAT ACT 309  
Cys Ser Asp Val Asn Asn Ser Thr Asn Pro Asn Asp Thr  
95 100

55 AAT ACT AAT TCC ACT AAT ACT ACT TCC TCT ACT CCT ACG 348  
Asn Thr Asn Ser Thr Asn Thr Thr Ser Ser Thr Pro Thr  
105 110 115

60 GCC ACT ACT AGT AGC GAG GAA AAG ATG GAG AAG GGA GAA 387  
Ala Thr Thr Ser Ser Glu Glu Lys Met Glu Lys Gly Glu  
120 125

65 ATA AAA AAC TGC TCT TTC AAT ATC ACC ACA CAC ATG AAA 426  
Ile Lys Asn Cys Ser Phe Asn Ile Thr Thr His Met Lys  
130 135 140

70 GAT AAG GTA CAG AAA GAA TAT GCA CTT TTT TAT AAA CTT 465  
Asp Lys Val Gin Lys Glu Tyr Ala Leu Phe Tyr Lys Leu  
145 150 155

GAT ATA GTA CCA ATA GAT GAT AAT AAT ACC AGC TAT AGG 504  
 Asp Ile Val Pro Ile Asp Asp Asn Asn Thr Ser Tyr Arg  
 160 165

5 TTG ATA AGT TGT AAT ACC TCA GTC ATT ACA CAG GCC TGT 543  
 Leu Ile Ser Cys Asn Thr Ser Val Ile Thr Gln Ala Cys  
 170 175 180

10 CCA ATG GTG ACC TTT GAG CCA ATT CCC ATA CAT TAT TGT 582  
 Pro Met Val Thr Phe Glu Pro Ile Pro Ile His Tyr Cys  
 185 190

15 GCC CCG GCT GGT TTT GCG ATT CTA AAG TGT AAA GAT AAG 621  
 Ala Pro Ala Gly Phe Ala Ile Leu Lys Cys Lys Asp Lys  
 195 200 205

20 AAG TTC AAT GGA ACA GGA CCA TGT TCA AAG GTC AGC ACA 660  
 Lys Phe Asn Gly Thr Gly Pro Cys Ser Lys Val Ser Thr  
 210 215 220

25 GTA CAA TGT ACA CAT GGA ATT ACG CCA GTA GTA TCA ACT 699  
 Val Gln Cys Thr His Gly Ile Arg Pro Val Val Ser Thr  
 225 230

30 CAA CTG TTG TTA AAT GGC AGT CTT GCA GAA GAA GAA GCA 738  
 Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu Glu Glu Val  
 235 240 245

35 GTA ATT AGA TCT GTC AAT TTC ACA CAC AAT CCT AAA ATC 777  
 Val Ile Arg Ser Val Asn Phe Thr Asp Asn Ala Lys Ile  
 250 255

40 ATA ATA GTA CAG CTG AAA GAA CCT GCA ATT AAT TGT 816  
 Ile Ile Val Gln Leu Lys Glu Pro Val Ala Ile Asn Cys  
 260 265 270

45 ACA AGA CCC AAC AAC AAT ACA AGA AAA CCT ATA CAT CTA 855  
 Thr Arg Pro Asn Asn Asn Thr Arg Lys Gly Ile His Leu  
 275 280 285

50 GGA CCA GGG AGC ACA TTT TAT ACA ACA GGA GAA ATA ATA 894  
 Gly Pro Gly Ser Thr Phe Tyr Thr Gly Glu Ile Ile  
 290 295

55 GGA GAC ATA AGA AAA GCA TAT TGC AAG ATT ACT AAA GAA 933  
 Gly Asp Ile Arg Lys Ala Tyr Cys Lys Ile Ser Lys Glu  
 300 305 310

60 AAA TGG AAT AAC ACT TTA AGA CAG GCA GTT AAA AAA TTA 972  
 Lys Trp Asn Asn Thr Leu Arg Gln Val Val Lys Lys Leu  
 315 320

65 AGA GAA CAA TTT GGG AAT AAA ACA ATA ATT TTT AAT CGA 1011  
 Arg Glu Gln Phe Gly Asn Lys Thr Ile Ile Phe Asn Arg  
 325 330 335

70 TCC TCA GGA CGG GAC CCA GAA ATT CTA ATG CAC AGT TTT 1050  
 Ser Ser Gly Gly Asp Pro Glu Ile Val Met His Ser Phe  
 340 345 350

75 AAC TGT GGA CGG GAG TTT TTC TAC TGT AAT ACA ACA CAA 1089  
 Asn Cys Gly Gly Glu Phe Phe Tyr Cys Asn Thr Thr Gln  
 355 360

1 CTG TTT AAT AGT ACT TGG AAT AAT ACT GAA GGG ACA AAT 1128  
 2 Leu Phe Asn Ser Thr Trp Asn Asn Thr Glu Gly Thr Asn  
 3 365 370 375  
 4 5 AGC ACT GAA GGA AAT AGC ACA ATC ACA CTC CCA TGC AGA 1167  
 5 Ser Thr Glu Gly Asn Ser Thr Ile Thr Leu Pro Cys Arg  
 6 380 385  
 7 10 ATA AAA CAA ATT ATA AAT ATG TGG CAG GAA GTC CGA AAA 1206  
 8 Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys  
 9 390 395 400  
 10 15 GCA ACG TAT GCC CCT CCC ATC AGA GGA CGA ATT AGA TGC 1245  
 11 Ala Thr Tyr Ala Pro Pro Ile Arg Gly Arg Ile Arg Cys  
 12 405 410 415  
 13 20 ATA TCA AAT ATT ACA GGA CTG CTA TTA ACA AGA GAT GGT 1284  
 14 Ile Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly  
 15 420 425  
 16 25 GGT AGG AAT GTC ACA AAC AAT ACC GAN NCC TTC AGA CCT 1323  
 17 Gly Arg Asn Val Thr Asn Asn Thr Xaa Xaa Phe Arg Pro  
 18 430 435 440  
 19 30 GGA GGA GGA GAC ATG AGG GAC AAT TGG AGA AGT GAA TTA 1362  
 20 Gly Gly Gly Asp Met Arg Asp Asn Trp Arg Ser Glu Leu  
 21 445 450  
 22 35 TAT AAA TAT AAA GTC GTC AAA GTT GAA CCA TTA GGA ATA 1401  
 23 Tyr Lys Tyr Lys Val Val Lys Val Glu Pro Leu Gly Ile  
 24 455 460 465  
 25 40 GCA CCC ACC AAG GCA AAC AGA AGA GTG GTG CAC AGA GAC 1440  
 26 Ala Pro Thr Lys Ala Lys Arg Arg Val Val His Arg Asp  
 27 470 475 480  
 28 45 (2) INFORMATION FOR SEQ ID NO:20:  
 29 (i) SEQUENCE CHARACTERISTICS:  
 30 (A) LENGTH: 499 amino acids  
 31 (B) TYPE: Amino Acid  
 32 (D) TOPOLOGY: Linear  
 33 50 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:20:  
 34 Glu Val Pro Val Trp Lys Glu Ala Thr Thr Leu Phe Cys Ala  
 35 1 5 10 15  
 36 55 Ser Asp Ala Lys Ala Tyr Asp Thr Glu Val His Asn Val Trp Ala  
 37 20 25 30  
 38 60 Thr His Ala Cys Val Pro Thr Asp Pro Asn Pro Gln Glu Ile Glu  
 39 35 40 45  
 40 65 Leu Val Asn Val Thr Glu Asp Phe Asn Met Trp Lys Asn Lys Met  
 41 50 55 60  
 42 65 Val Asp Gln Met His Glu Asp Ile Ile Ser Leu Trp Asp Glu Ser  
 43 70 75

	Leu	Lys	Pro	Cys	Val	Lys	Leu	Thr	Pro	Leu	Cys	Val	Thr	Leu	Asn
															80
															85
															90
5	Cys	Ser	Asp	Val	Asn	Asn	Ser	Thr	Asn	Pro	Asn	Asp	Thr	Asn	Thr
															95
															100
															105
	Asn	Ser	Thr	Asn	Thr	Thr	Ser	Ser	Thr	Pro	Thr	Ala	Thr	Thr	Ser
															110
															115
															120
10	Ser	Glu	Glu	Lys	Met	Glu	Lys	Gly	Glu	Ile	Lys	Asn	Cys	Ser	Phe
															125
															130
															135
	Asn	Ile	Thr	Thr	His	Met	Lys	Asp	Lys	Val	Gln	Lys	Glu	Tyr	Ala
															140
															145
15	Leu	Phe	Tyr	Lys	Leu	Asp	Ile	Val	Pro	Ile	Asp	Asp	Asn	Asn	Thr
															155
															160
															165
20	Ser	Tyr	Arg	Leu	Ile	Ser	Cys	Asn	Thr	Ser	Val	Ile	Thr	Gln	Ala
															170
															175
															180
	Cys	Pro	Met	Val	Thr	Phe	Glu	Pro	Ile	Pro	Ile	His	Tyr	Cys	Ala
															185
															190
															195
25	Pro	Ala	Gly	Phe	Ala	Ile	Leu	Lys	Cys	Lys	Asp	Lys	Lys	Phe	Asn
															200
															205
															210
	Gly	Thr	Gly	Pro	Cys	Ser	Lys	Val	Ser	Thr	Val	Gln	Cys	Thr	His
															215
															220
30	Gly	Ile	Arg	Pro	Val	Val	Ser	Thr	Gln	Leu	Leu	Leu	Asn	Gly	Ser
															230
															235
															240
35	Leu	Ala	Glu	Glu	Glu	Val	Val	Ile	Arg	Ser	Val	Asn	Phe	Thr	Asp
															245
															250
															255
	Asn	Ala	Lys	Ile	Ile	Ile	Val	Gln	Leu	Lys	Glu	Pro	Val	Ala	Ile
															260
															265
40	Asn	Cys	Thr	Arg	Pro	Asn	Asn	Asn	Thr	Arg	Lys	Gly	Ile	His	Leu
															275
															280
															285
	Gly	Pro	Gly	Ser	Thr	Phe	Tyr	Thr	Thr	Gly	Glu	Ile	Ile	Gly	Asp
															290
															295
45	Ile	Arg	Lys	Ala	Tyr	Cys	Lys	Ile	Ser	Lys	Glu	Lys	Trp	Asn	Asn
															305
															310
															315
50	Thr	Leu	Arg	Gln	Val	Val	Lys	Lys	Leu	Arg	Glu	Gln	Phe	Gly	Asn
															320
															325
															330
	Lys	Thr	Ile	Ile	Phe	Asn	Arg	Ser	Ser	Gly	Gly	Asp	Pro	Glu	Ile
															335
															340
55	Val	Met	His	Ser	Phe	Asn	Cys	Gly	Gly	Glu	Phe	Phe	Tyr	Cys	Asn
															350
															355
															360
	Thr	Thr	Gln	Leu	Phe	Asn	Ser	Thr	Trp	Asn	Asn	Thr	Glu	Gly	Thr
															365
															370
60	Asn	Ser	Thr	Glu	Gly	Asn	Ser	Thr	Ile	Thr	Leu	Pro	Cys	Arg	Ile
															380
															385
	Lys	Gln	Ile	Ile	Asn	Met	Trp	Gln	Glu	Val	Gly	Lys	Ala	Thr	Tyr
65															395
															400
															405

Ala Pro Pro Ile Arg Gly Arg Ile Arg Cys Ile Ser Asn Ile Thr  
 410 415 420  
 Gly Leu Leu Leu Thr Arg Asp Gly Gly Arg Asn Val Thr Asn Asn  
 5 425 430 435  
 Thr Xaa Xaa Phe Arg Pro Gly Gly Asp Met Arg Asp Asn Trp  
 440 445 450  
 10 Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Val Glu Pro Leu  
 455 460 465  
 Gly Ile Ala Pro Thr Lys Ala Lys Arg Arg Val Val His Arg Asp  
 470 475 480  
 15 Lys Arg Ala Ala Leu Gly Ala Leu Phe Leu Gly Phe Leu Gly Ala  
 485 490 495  
 Xaa Lys Leu Leu  
 20 499

## (2) INFORMATION FOR SEQ ID NO:21:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 1475 base pairs
- (B) TYPE: Nucleic Acid
- (C) STRANDEDNESS: Single
- (D) TOPOLOGY: Linear

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:21:

30 G GTA CCT GTG TGG AAA GAA GCA AAC ACA ACT CTA TTT 37  
 Val Pro Val Trp Lys Glu Ala Asn Thr Thr Leu Phe  
 1 5 10  
 TGT GCA TCA GAT GCT AAA GCA TAT GAT AGA GAA GTA CAT 76  
 Cys Ala Ser Asp Ala Lys Ala Tyr Asp Arg Glu Val His  
 35 15 20 25  
 AAT GTT TGG GCA ACA CAT GCC TGT GTC CCC ACA GAC CCC 115  
 Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro  
 40 30 35  
 AAC CCA CAA GAA ATA GTA TTG GGA AAT GTG ACA GAA AAT 154  
 Asn Pro Gln Glu Ile Val Leu Gly Asn Val Thr Glu Asn  
 45 40 45 50  
 TTT AAC ATG TGG AAA AAT AAC ATG GTC GAA CAA ATG CAT 193  
 Phe Asn Met Trp Lys Asn Asn Met Val Glu Gln Met His  
 55 55 60  
 50 GAG GAT ATA ATC AAT TTA TGG GAT CAA AGC TTA AAG CCA 232  
 Glu Asp Ile Ile Asn Leu Trp Asp Gln Ser Leu Lys Pro  
 65 65 70 75  
 TGT GTC AAG TTA ACT CCA CTC TGT GTT ACT TTA AAG TGC 271  
 Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu Lys Cys  
 55 80 85 90  
 AAG GAT CTG GAG AGG AAT ACT ACC TAT AAT AGC ACT ATT 310  
 Lys Asp Leu Glu Arg Asn Thr Thr Tyr Asn Ser Thr Ile  
 60 95 100  
 ACC AAT AAT AGT AGT TTG GAG GGA CTA AGA GAA CAA ATG 349  
 Thr Asn Asn Ser Ser Leu Glu Gly Leu Arg Glu Gln Met  
 105 110 115  
 65

ACA AAC TGC TCT TTC AAC ATC ACC ACA AGT ATA AGA GAT 388  
 Thr Asn Cys Ser Phe Asn Ile Thr Thr Ser Ile Arg Asp  
 120 125

5 AAG CTG CAG AAA GAA TAT GCA CTT TTG TAT AAA CTT GAT 427  
 Lys Val Gln Lys Glu Tyr Ala Leu Leu Tyr Lys Leu Asp  
 130 135 140

10 GTA CTA CCA ATA GAA GAA GAT GAC AAT ACT AGC TAT AGA 466  
 Val Val Pro Ile Glu Glu Asp Asp Asn Thr Ser Tyr Arg  
 145 150 155

TTG ATA AGT TGT AAC ACC TCA GTC ATT ACA CAG GCT TGT 505  
 Leu Ile Ser Cys Asn Thr Ser Val Ile Thr Gln Ala Cys  
 15 160 165

CCA AAG ACA TCC TTT GAG CCA ATT CCC ATA CAT TAT TGT 544  
 Pro Lys Thr Ser Phe Glu Pro Ile Pro Ile His Tyr Cys  
 170 175 180

20 GCC CCG GCT GGT TTT GCG ATT CTA AAG TGT AAT GAT AAG 583  
 Ala Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asp Lys  
 185 190

25 AAG TTC AAT GGA ACA GGA CCA TGT AAA AAT CTC ACC ACA 622  
 Lys Phe Asn Gly Thr Gly Pro Cys Lys Asn Val Ser Thr  
 195 200 205

30 GTA CAA TGT ACA CAT GGA ATT AGG CCA GTA GTA TCA ACT 661  
 Val Gln Cys Thr His Gly Ile Arg Pro Val Val Ser Thr  
 210 215 220

CAA CTG TTG TTA AAT GGC AGT CTA GCA GAA GAA GAG GTA 700  
 Gln Leu Leu Asn Gly Ser Leu Ala Glu Glu Glu Val  
 35 225 230

GTA ATC AGA TCT GCC AAT TTC ACA GAC AAT GCT AAA ACC 739  
 Val Ile Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr  
 235 240 245

40 ATA ATA GTA CAT CTA AAT GAA ACT GTA AAA ATT AAT TGT 778  
 Ile Ile Val His Leu Asn Glu Thr Val Lys Ile Asn Cys  
 250 255

45 ACA AGA CTT GGC AAC AAT ACA AGA AAA AGT ATA AAT ATA 817  
 Thr Arg Leu Gly Asn Asn Thr Arg Lys Ser Ile Asn Ile  
 260 265 270

50 GGA CCA CGG AGA GTA CTC TAT GCA ACA GGA GAA ATA ATA 856  
 Gly Pro Gly Arg Val Leu Tyr Ala Thr Gly Glu Ile Ile  
 275 280 285

55 GGA GAC ATA AGA CAA GCA CAT TGT AAC ATT AGT AGA GCA 895  
 Gly Asp Ile Arg Gln Ala His Cys Asn Ile Ser Arg Ala  
 290 295

CAA TGG AAT AAG ACT TTA GAA AAG CTA GTT GAC AAA TTA 934  
 Gln Trp Asn Lys Thr Leu Glu Lys Val Val Asp Lys Leu  
 300 305 310

60 AGA AAA CAA TTT GGG GAT AAT ACA ACA ATA GCT TTT AAT 973  
 Arg Lys Gln Phe Gly Asp Asn Thr Thr Ile Ala Phe Asn  
 315 320

CGA TCC TCA GGA GGG GAC CCA GAA ATT GTA ATG CAC ACT 1012  
 Arg Ser Ser Gly Gly Asp Pro Glu Ile Val Met His Thr  
 325 330 335  
 5 TTT AAT TCT GGA GGG GAA TTT TTC TAC TGT AAT ACA ACA 1051  
 Phe Asn Cys Gly Gly Glu Phe Phe Tyr Cys Asn Thr Thr  
 340 345 350  
 10 CAA CTG TTT AAT AGT ACT TGG AAT AAT ACT TGG AAG GAT 1090  
 Gln Leu Phe Asn Ser Thr Trp Asn Asn Thr Trp Lys Asp  
 355 360  
 15 CCT AAC AGG AGT GAC AAT ATC ACA CTC CCA TGC AGA ATA 1129  
 Pro Asn Arg Ser Asp Asn Ile Thr Leu Pro Cys Arg Ile  
 365 370 375  
 20 AAA CAA ATT ATA AAC ATG TGG CAG GAA GTA GGA AAA GCA 1168  
 Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys Ala  
 380 385  
 25 ATG TAC GCC CCT CCC ATC AGA GGG GAA ATT AGA TGT TCA 1207  
 Met Tyr Ala Pro Pro Ile Arg Gly Glu Ile Arg Cys Ser  
 390 395 400  
 30 TCA AAT ATC ACA GGG CTG CTA CTA ACA AGA GAT GGT GGT 1246  
 Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly  
 405 410 415  
 AAT GAC GAT GGT AAT GAC ACG ACC ACA AAC AGG ACC GAG 1285  
 35 Asn Asp Asp Gly Asn Asp Thr Thr Thr Asn Arg Thr Glu  
 420 425  
 ATC TTC AGA CCT GGA GGA GAT ATG AGG GAC AAT TGG 1324  
 Ile Phe Arg Pro Gly Gly Asp Met Arg Asp Asn Trp  
 430 435 440  
 40 AGA AGT GAA TTA TAT AGA TAT AAA GTA GTA AAA ATT GAA 1363  
 Arg Ser Glu Leu Tyr Arg Tyr Lys Val Val Lys Ile Glu  
 445 450  
 45 CCA TTA GGA ATA GCA CCC ACC AGG GCA AAG AGA AGA GTG 1402  
 Pro Leu Gly Ile Ala Pro Thr Arg Ala Lys Arg Arg Val  
 455 460 465  
 GTG CAG AGA GAA AAA AGA GCA GTA GGA CTA GGA GCT TTG 1441  
 Val Gln Arg Glu Lys Arg Ala Val Gly Leu Gly Ala Leu  
 470 475 480  
 50 TTC CTT GGG T TCTTAGGAG CATAAGCTT CTAGA 1475  
 Phe Leu Gly  
 483

## (2) INFORMATION FOR SEQ ID NO:22:

## (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 491 amino acids

(B) TYPE: Amino Acid

(D) TOPOLOGY: Linear

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:22:

60 Val Pro Val Trp Lys Glu Ala Asn Thr Thr Leu Phe Cys Ala Ser  
 1 5 10 15  
 Asp Ala Lys Ala Tyr Asp Arg Glu Val His Asn Val Trp Ala Thr  
 20 25 30  
 65

	His	Ala	Cys	Val	Pro	Thr	Asp	Pro	Asn	Pro	Gln	Glu	Ile	Val	Leu
35															45
5	Gly	Asn	Val	Thr	Glu	Asn	Phe	Asn	Met	Trp	Lys	Asn	Asn	Met	Val
														60	
	Glu	Gln	Met	His	Glu	Asp	Ile	Ile	Asn	Leu	Trp	Asp	Gln	Ser	Leu
															75
10	Lys	Pro	Cys	Val	Lys	Leu	Thr	Pro	Leu	Cys	Val	Thr	Leu	Lys	Cys
															90
	Lys	Asp	Leu	Glu	Arg	Asn	Thr	Thr	Tyr	Asn	Ser	Thr	Ile	Thr	Asn
															105
15	Asn	Ser	Ser	Leu	Glu	Gly	Leu	Arg	Glu	Gln	Met	Thr	Asn	Cys	Ser
															120
20	Phe	Asn	Ile	Thr	Thr	Ser	Ile	Arg	Asp	Lys	Val	Gln	Lys	Glu	Tyr
															135
	Ala	Leu	Leu	Tyr	Lys	Leu	Asp	Val	Val	Pro	Ile	Glu	Glu	Asp	Asp
															150
25	Asn	Thr	Ser	Tyr	Arg	Leu	Ile	Ser	Cys	Asn	Thr	Ser	Val	Ile	Thr
															165
	Gln	Ala	Cys	Pro	Lys	Thr	Ser	Phe	Glu	Pro	Ile	Pro	Ile	His	Tyr
															180
30	Cys	Ala	Pro	Ala	Gly	Phe	Ala	Ile	Leu	Lys	Cys	Asn	Asp	Lys	Lys
															195
35	Phe	Asn	Gly	Thr	Gly	Pro	Cys	Lys	Asn	Val	Ser	Thr	Val	Gln	Cys
															210
	Thr	His	Gly	Ile	Arg	Pro	Val	Val	Ser	Thr	Gln	Leu	Leu	Leu	Asn
															225
40	Gly	Ser	Ile	Ala	Glu	Glu	Val	Val	Ile	Arg	Ser	Ala	Asn	Phe	
															240
	Thr	Asp	Asn	Ala	Lys	Thr	Ile	Ile	Val	His	Leu	Asn	Glu	Thr	Val
															255
45	Lys	Ile	Asn	Cys	Thr	Arg	Leu	Gly	Asn	Asn	Thr	Arg	Lys	Ser	Ile
															270
50	Asn	Ile	Gly	Pro	Gly	Arg	Val	Leu	Tyr	Ala	Thr	Gly	Glu	Ile	Ile
															285
	Gly	Asp	Ile	Arg	Gln	Ala	His	Cys	Asn	Ile	Ser	Arg	Ala	Gln	Trp
															300
55	Asn	Lys	Thr	Leu	Glu	Lys	Val	Val	Asp	Lys	Leu	Arg	Lys	Gln	Phe
															315
60	Gly	Asp	Asn	Thr	Thr	Ile	Ala	Phe	Asn	Arg	Ser	Ser	Gly	Gly	Asp
															330
	Pro	Glu	Ile	Val	Met	His	Thr	Phe	Asn	Cys	Gly	Gly	Glu	Phe	Phe
															345
65	Tyr	Cys	Asn	Thr	Thr	Gln	Leu	Phe	Asn	Ser	Thr	Trp	Asn	Asn	Thr
															360

Trp Lys Asp Pro Asn Arg Ser Asp Asn Ile Thr Leu Pro Cys Arg  
 365 370 375  
 Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys Ala Met  
 5 380 385 390  
 Tyr Ala Pro Pro Ile Arg Gly Glu Ile Arg Cys Ser Ser Asn Ile  
 395 400 405  
 10 Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly Asn Asp Asp Gly Asn  
 410 415 420  
 Asp Thr Thr Thr Asn Arg Thr Glu Ile Phe Arg Pro Gly Gly Gly  
 425 430 435  
 15 Asp Met Arg Asp Asn Trp Arg Ser Glu Leu Tyr Arg Tyr Lys Val  
 440 445 450  
 20 Val Lys Ile Glu Pro Leu Gly Ile Ala Pro Thr Arg Ala Lys Arg  
 455 460 465  
 Arg Val Val Gln Arg Glu Lys Arg Ala Val Gly Leu Gly Ala Leu  
 470 475 480  
 25 Phe Leu Gly Phe Leu Gly Ala Leu Phe Leu Gly  
 485 490 491

## (2) INFORMATION FOR SEQ ID NO:23:

## (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1475 base pairs

(B) TYPE: Nucleic Acid

(C) STRANDEDNESS: Single

(D) TOPOLOGY: Linear

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:23:

35 G GTA CCT GTG TGG AAA GAA GCA AAC ACA ACT CTA TTT 37  
 Val Pro Val Trp Lys Glu Ala Asn Thr Thr Leu Phe  
 1 5 10  
 40 TGT GCA TCA GAT GCT AAA GCA TAT GAT AGA GAA GTA CAT 76  
 Cys Ala Ser Asp Ala Lys Ala Tyr Asp Arg Glu Val His  
 15 20 25  
 45 AAT GTT TGG GCA ACA CAT GCC TGT GTA CCC ACA GAC CCC 115  
 Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro  
 30 35  
 50 AAC CCA CAA GAA ATA GTA TTG GGA AAT GTG ACA GAA AAT 154  
 Asn Pro Gln Glu Ile Val Leu Gly Asn Val Thr Glu Asn  
 40 45 50  
 55 TTT AAC ATG TGG AAA AAT AAC ATG GTA GAA CAA ATG CAT 193  
 Phe Asn Met Trp Lys Asn Asn Met Val Glu Gln Met His  
 55 60  
 60 GAG GAT ATA ATC AAT TTA TGG GAT CAA AGC TTA AAG CCA 232  
 Glu Asp Ile Ile Asn Leu Trp Asp Gln Ser Leu Lys Pro  
 65 70 75  
 60 TGT GTA AAG TTA ACT CCA CTC TGT GTT ACT TTA AAG TGC 271  
 Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu Lys Cys  
 80 85 90

1 AAG GAT CTG GAG AGG AAT ACT ACC TAT AAT AGC ACT ATT 310  
 Lys Asp Leu Glu Arg Asn Thr Thr Tyr Asn Ser Thr Ile  
 95 100

5 ACC AAT AAT AGT AGT TTG GAG CGA CTA AGA GAA CAA ATG 349  
 Thr Asn Asn Ser Ser Leu Glu Gly Leu Arg Glu Gln Met  
 105 110 115

10 ACA AAC TGC TCT TTC AAC ATC ACC ACA AGT ATA AGA GAT 388  
 Thr Asn Cys Ser Phe Asn Ile Thr Thr Ser Ile Arg Asp  
 120 125

15 AAG CTG CAG AAA GAA TAT GCA CTT TTG TAT AAA CTT GAT 427  
 Lys Val Gln Lys Glu Tyr Ala Leu Leu Tyr Lys Leu Asp  
 130 135 140

20 GTA GTA CCA ATA GAA GAA GAT GAC AAT ACT AGC TAT AGA 466  
 Val Val Pro Ile Glu Glu Asp Asp Asn Thr Ser Tyr Arg  
 145 150 155

25 TTG ATA AGT TGT AAC ACC TCA GTC ATT ACA CAG GCT TGT 505  
 Leu Ile Ser Cys Asn Thr Ser Val Ile Thr Gln Ala Cys  
 160 165

30 CCA AAG ACA TCC TTT GAG CCA ATT CCC ATA CAT TAT TGT 544  
 Pro Lys Thr Ser Phe Glu Pro Ile Pro Ile His Tyr Cys  
 170 175 180

35 CCC CCG GCT GGT TTT GCG ATT CTA AAG TGT AAT GAT AAG 583  
 Ala Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asp Lys  
 185 190

40 AAG TTC AAT GGA ACA GGA CCA TGT AAA AAT GTC AGC ACA 622  
 Lys Phe Asn Gly Thr Gly Pro Cys Lys Asn Val Ser Thr  
 195 200 205

45 GTA CAA TGT ACA CAT GCA ATT AGG CCA GTA GTA TCA ACT 661  
 Val Gln Cys Thr His Gly Ile Arg Pro Val Val Ser Thr  
 210 215 220

50 CAA CTG TTG TTA AAT GGC AGT CTA GCA GAA GAA GAG GTA 700  
 Gln Leu Leu Asn Gly Ser Leu Ala Glu Glu Glu Val  
 225 230

55 ATA ATA GTA CAT CTA AAT GAA ACT GTA AAA ATT AAT TGT 778  
 Ile Ile Val His Leu Asn Glu Thr Val Lys Ile Asn Cys  
 235 240 245

60 ACA AGA CTT GGC AAC AAT ACA AGA AAA AGT ATA AAT ATA 817  
 Thr Arg Leu Gly Asn Asn Thr Arg Lys Ser Ile Asn Ile  
 260 265 270

65 GGA CCA GGG AGA GTA CTC TAT GCA ACA GGA GAA ATA ATA 856  
 Gly Pro Gly Arg Val Leu Tyr Ala Thr Gly Glu Ile Ile  
 275 280 285

70 GGA GAC ATA AGA CAA GCA CAT TGT AAC ATT AGT AGA GCA 895  
 Gly Asp Ile Arg Gln Ala His Cys Asn Ile Ser Arg Ala  
 290 295

CAA TGG AAT AAG ACT TTA GAA AAG GTA GTT GAC AAG TTA 934  
 Gln Trp Asn Lys Thr Leu Glu Lys Val Val Asp Lys Leu  
 300 305 310  
 5 AGA AAA CAA TTT GGG GAT AAT ACA ACA ATA GCT TTT AAT 973  
 Arg Lys Gln Phe Gly Asp Asn Thr Thr Ile Ala Phe Asn  
 315 320  
 CGA TCC TCA GGA GGG GAC CCA GAA ATT GTA ATG CAC ACT 1012  
 10 Arg Ser Ser Gly Gly Asp Pro Glu Ile Val Met His Thr  
 325 330 335  
 TTT AAT TGT GGA GGG GAA TTT TTC TAC TGT AAT ACA ACA 1051  
 Phe Asn Cys Gly Gly Glu Phe Phe Tyr Cys Asn Thr Thr  
 15 340 345 350  
 CAA CTG TTT AAT AGT ACT TGG AAT AAT ACT TGG AAG GAT 1090  
 Gln Leu Phe Asn Ser Thr Trp Asn Asn Thr Trp Lys Asp  
 355 360  
 20 CCT AAC AGG AGT GAC AAT ATC ACA CTC CCA TGC AGA ATA 1129  
 Pro Asn Arg Ser Asp Asn Ile Thr Leu Pro Cys Arg Ile  
 365 370 375  
 25 AAA CAA ATT ATA AAC ATG TGG CAG GAA GTA GGA AAA GCA 1168  
 Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys Ala  
 380 385  
 ATG TAC GCC CCT CCC ATC AGA GGG GAA ATT AGA TGT TCA 1207  
 30 Met Tyr Ala Pro Pro Ile Arg Gly Glu Ile Arg Cys Ser  
 390 395 400  
 TCA AAT ATC ACA GGG CTG CTA CTA ACA AGA GAT GGT GGT 1246  
 Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly  
 35 405 410 415  
 AAT GAC GAT GGT AAT GAC ACG ACC ACA AAC AGG ACC GAG 1285  
 Asn Asp Asp Gly Asn Asp Thr Thr Asn Arg Thr Glu  
 420 425  
 40 ATC TTC AGA CCT GGA GGA GGA GAT ATG AGG GAC AAT TGG 1324  
 Ile Phe Arg Pro Gly Gly Asp Met Arg Asp Asn Trp  
 430 435 440  
 45 AGA AGT GAA TTA TAT AGA TAT AAA GTA GTA AAA ATT GAA 1363  
 Arg Ser Glu Leu Tyr Arg Tyr Lys Val Val Lys Ile Glu  
 445 450  
 CCA TTA GGA ATA GCA CCC ACC AGG GCA AAG AGA AGA GTG 1402  
 50 Pro Leu Gly Ile Ala Pro Thr Arg Ala Lys Arg Arg Val  
 455 460 465  
 GTG CAG AGA GAA AAA AGA GCA GTA GGA CTA GGA GCT TTG 1441  
 Val Gln Arg Glu Lys Arg Ala Val Gly Leu Gly Ala Leu  
 55 470 475 480  
 TTC CTT GGG TTC TTG GGA CCA TAA AGC TTC TAG A 1475  
 Phe Leu Gly Phe Leu Gly Ala Xaa Ser Phe Xaa  
 485 490 491  
 60 (2) INFORMATION FOR SEQ ID NO:24:  
 (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 491 amino acids  
 (B) TYPE: Amino Acid  
 (C) TOPOLOGY: Linear

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:24:

1	Val	Pro	Val	Trp	Lys	Glu	Ala	Asn	Thr	Thr	Leu	Phe	Cys	Ala	Ser
5	1	5							10				15		
5	Asp	Ala	Lys	Ala	Tyr	Asp	Arg	Glu	Val	His	Asn	Val	Trp	Ala	Thr
		20						25					30		
10	His	Ala	Cys	Val	Pro	Thr	Asp	Pro	Asn	Pro	Gln	Gl	Ile	Val	Leu
		35						40				45			
	Gly	Asn	Val	Thr	Glu	Asn	Phe	Asn	Met	Trp	Lys	Asn	Asn	Met	Val
		50			55							60			
15	Glu	Gln	Met	His	Glu	Asp	Ile	Ile	Asn	Leu	Trp	Asp	Gln	Ser	Leu
		65						70				75			
	Lys	Pro	Cys	Val	Lys	Leu	Thr	Pro	Leu	Cys	Val	Thr	Leu	Lys	Cys
		80						85				90			
20	Lys	Asp	Leu	Glu	Arg	Asn	Thr	Thr	Tyr	Asn	Ser	Thr	Ile	Thr	Asn
		95						100				105			
25	Asn	Ser	Ser	Leu	Glu	Gly	Leu	Arg	Glu	Gln	Met	Thr	Asn	Cys	Ser
		110						115				120			
	Phe	Asn	Ile	Thr	Thr	Ser	Ile	Arg	Asp	Lys	Val	Gln	Lys	Glu	Tyr
		125						130				135			
30	Ala	Leu	Leu	Tyr	Lys	Leu	Asp	Val	Val	Pro	Ile	Glu	Glu	Asp	Asp
		140						145				150			
	Asn	Thr	Ser	Tyr	Arg	Leu	Ile	Ser	Cys	Asn	Thr	Ser	Val	Ile	Thr
		155						160				165			
35	Gln	Ala	Cys	Pro	Lys	Thr	Ser	Phe	Glu	Pro	Ile	Pro	Ile	His	Tyr
		170						175				180			
40	Cys	Ala	Pro	Ala	Gly	Phe	Ala	Ile	Leu	Lys	Cys	Asn	Asp	Lys	Lys
		185						190				195			
	Phe	Asn	Gly	Thr	Gly	Pro	Cys	Lys	Asn	Val	Ser	Thr	Val	Gln	Cys
		200						205				210			
45	Thr	His	Gly	Ile	Arg	Pro	Val	Val	Ser	Thr	Gln	Leu	Leu	Leu	Asn
		215						220				225			
	Gly	Ser	Leu	Ala	Glu	Glu	Val	Val	Ile	Arg	Ser	Ala	Asn	Phe	
		230						235				240			
50	Thr	Asp	Asn	Ala	Lys	Thr	Ile	Ile	Val	His	Leu	Asn	Glu	Thr	Val
		245						250				255			
	Lys	Ile	Asn	Cys	Thr	Arg	Leu	Gly	Asn	Asn	Thr	Arg	Lys	Ser	Ile
		260						265				270			
55	Asn	Ile	Gly	Pro	Gly	Arg	Val	Leu	Tyr	Ala	Thr	Gly	Glu	Ile	Ile
		275						280				285			
60	Gly	Asp	Ile	Arg	Gln	Ala	His	Cys	Asn	Ile	Ser	Arg	Ala	Gln	Trp
		290						295				300			
	Asn	Lys	Thr	Leu	Glu	Lys	Val	Val	Asp	Lys	Leu	Arg	Lys	Gln	Phe
		305						310				315			
65															

Gly Asp Asn Thr Thr Ile Ala Phe Asn Arg Ser Ser Gly Gly Asp  
 320 325 330  
 Pro Glu Ile Val Met His Thr Phe Asn Cys Gly Gly Glu Phe Phe  
 5 335 340 345  
 Tyr Cys Asn Thr Thr Gln Leu Phe Asn Ser Thr Trp Asn Asn Thr  
 350 355 360  
 10 Trp Lys Asp Pro Asn Arg Ser Asp Asn Ile Thr Leu Pro Cys Arg  
 365 370 375  
 Ile Lys Gin Ile Ile Asn Met Trp Gln Glu Val Gly Lys Ala Met  
 380 385 390  
 15 Tyr Ala Pro Pro Ile Arg Gly Glu Ile Arg Cys Ser Ser Asn Ile  
 395 400 405  
 Thr Gly Leu Leu Leu Thr Arg Asp Gly Asn Asp Asp Gly Asn  
 20 410 415 420  
 Asp Thr Thr Thr Asn Arg Thr Glu Ile Phe Arg Pro Gly Gly  
 425 430 435  
 25 Asp Met Arg Asp Asn Trp Arg Ser Glu Leu Tyr Arg Tyr Lys Val  
 440 445 450  
 Val Lys Ile Glu Pro Leu Gly Ile Ala Pro Thr Arg Ala Lys Arg  
 455 460 465  
 30 Arg Val Val Gln Arg Glu Lys Arg Ala Val Gly Leu Gly Ala Leu  
 470 475 480  
 Phe Leu Gly Phe Leu Gly Ala Xaa Ser Phe Xaa  
 35 485 490 491

(2) INFORMATION FOR SEQ ID NO:25:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1435 base pairs

(B) TYPE: Nucleic Acid

(C) STRANDEDNESS: Single

(D) TOPOLOGY: Linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:25:

45 CTC GAG CCT GTG TGG AAA GAA GCA ACC ACC ACT 36  
 Leu Glu Val Pro Val Trp Lys Glu Ala Thr Thr Thr  
 1 5 10

50 CTA TTT TGT GCA TCA GAT GCT AAA GCA TAT GAT TCA GAG 75  
 Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Ser Glu  
 15 20 25

55 GCA CAT AAT GTT TGG GCC ACA CAT GCC TGT GTA CCC ACA 114  
 Ala His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr  
 30 35

60 GAC CCC AAC CCA CAA GAA GTA GAA TTG GAA AAT GTG ACA 153  
 Asp Pro Asn Pro Gln Glu Val Glu Leu Glu Asn Val Thr  
 40 45 50

GAA AAT TTT AAC ATG TGG AAA AAT AAC ATG GTA GAA CAG 192  
 Glu Asn Phe Asn Met Trp Lys Asn Asn Met Val Glu Gln  
 55 60

ATG CAT GGG GAT ATA ATT AGT TTA TGG GAT CAA AGC CTA 231  
 Met His Gly Asp Ile Ile Ser Leu Trp Asp Gln Ser Leu  
 65 70 75  
 5 AAG CCA TGT GTA AAA TTA ACC CCA CTC TGT GTT ACG TTA 270  
 Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu  
 80 85 90  
 10 AAT TGC ACT GAC CCA AAT GTT ACT AAT AGC GAG AGA ACG 309  
 Asn Cys Thr Asp Pro Asn Val Thr Asn Ser Glu Arg Thr  
 95 100  
 15 ATA GAG GGG GGA GAA ATA AAA AAT TGC TCT TTC AAT ATC 348  
 Ile Glu Gly Glu Ile Lys Asn Cys Ser Phe Asn Ile  
 105 110 115  
 20 ACC ACA AAC ATA AGA GAT AGG TTT CAG AAA GAA TAT GCA 387  
 Thr Thr Asn Ile Arg Asp Arg Phe Gln Lys Glu Tyr Ala  
 120 125  
 25 CTT TTT TAT AAA CTT GAT GTA ATA CCA TTA GGT AAT GAT 426  
 Leu Phe Tyr Lys Leu Asp Val Ile Pro Leu Gly Asn Asp  
 130 135 140  
 30 AAT ACT AGC TAT AGG TTG ATA AGT TGT AAC ACC TCA GTC 465  
 Asn Thr Ser Tyr Arg Leu Ile Ser Cys Asn Thr Ser Val  
 145 150 155  
 35 ATT ACA CAG GCC TGT CCA AAG GTA TCC TTT GAG CCA ATT 504  
 Ile Thr Gln Ala Cys Pro Lys Val Ser Phe Glu Pro Ile  
 160 165  
 CCC ATA CAT TAT TGT GCC CCG GCT GGT TTT CCG ATT CTA 543  
 Pro Ile His Tyr Cys Ala Pro Ala Gly Phe Ala Ile Leu  
 35 170 175 180  
 40 AAG TGT AAA GAT AAG AAG TTC AAT GGA ACA GGA CCA TGT 582  
 Lys Cys Lys Asp Lys Lys Phe Asn Gly Thr Gly Pro Cys  
 185 190  
 45 ACA AAT GTC AGC ACA GTA CAA TGT ACA CAT GGA ATT AAG 621  
 Thr Asn Val Ser Thr Val Gln Cys Thr His Gly Ile Lys  
 195 200 205  
 50 CCA GTA GTA TCA ACT CAA CTG TTG TTA AAT GGC ACT CTA 660  
 Pro Val Val Ser Thr Gln Leu Leu Leu Asn Gly Ser Leu  
 210 215 220  
 55 GCA GAA GAA GAC ATA GTA ATT AGA TCC GCC AAT CTC ACA 699  
 Ala Glu Glu Asp Ile Val Ile Arg Ser Ala Asn Leu Thr  
 225 230  
 60 GAC AAT GCT AAA AAC ATA ATA GTA CAG CTG AAT GAA TCT 738  
 Asp Asn Ala Lys Asn Ile Ile Val Gln Leu Asn Glu Ser  
 235 240 245  
 65 GTA ACA ATG AAT TGT ACA AGA CCC AAC AAC AAT ACA ATG 777  
 Val Thr Met Asn Cys Thr Arg Pro Asn Asn Asn Thr Met  
 250 255  
 70 AAA AGT ATA CAT ATA GGA CCA GCC AGA GCA TTT TAT GCA 816  
 Lys Ser Ile His Ile Gly Pro Gly Arg Ala Phe Tyr Ala  
 260 265 270

ACA GGA AAC ATA ATA GGA GAT ATA AGA CAA GCA CAT TGT 855  
 Thr Gly Asn Ile Ile Gly Asp Ile Arg Gln Ala His Cys  
 275 280 285  
 AAC ATT AGT GGA ACA AAA TGG AAT GAC ACT TTG AAA AAG 894  
 Asn Ile Ser Gly Thr Lys Trp Asn Asp Thr Leu Lys Lys  
 290 295  
 ATA GCT ATA AAA TTA AGA GAA CAA TTT AAT AAG ACA ATA 933  
 Ile Ala Ile Lys Leu Arg Glu Gln Phe Asn Lys Thr Ile  
 300 305 310  
 GTC TTT AAT CAA TCC TCA GGA GGG GAC CCA GAA ATT GCA 972  
 Val Phe Asn Gln Ser Ser Gly Gly Asp Pro Glu Ile Ala  
 315 320  
 ACG CTC AGT TTT AAT TGT GGA GGG GAA TTT TTC TAC TGT 1011  
 Thr Leu Ser Phe Asn Cys Gly Gly Glu Phe Phe Tyr Cys  
 325 330 335  
 AAT TCA ACA CAA CTG TTT AAT AGT ACT TGG AAT AGT ACT 1050  
 Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp Asn Ser Thr  
 340 345 350  
 GGG TCA AAT AAC ACT AAA GGA AAT GAC ACA ATC ACA CTC 1089  
 Gly Ser Asn Asn Thr Lys Gly Asn Asp Thr Ile Thr Leu  
 355 360  
 CCA TGC AGA ATA AGA CAA ATT ATA AAC ATG TGG CAG AAA 1128  
 Pro Cys Arg Ile Arg Gln Ile Ile Asn Met Trp Gln Lys  
 365 370 375  
 ATA GGA AAA GCA ATG TAT GCC CCT CCC ATC AAA GGG CAA 1167  
 Ile Gly Lys Ala Met Tyr Ala Pro Pro Ile Lys Gly Gln  
 380 385  
 ATT AGA TGT TCA TCA AAT ATT ACA GGG CTA ATA TTA ACA 1206  
 Ile Arg Cys Ser Ser Asn Ile Thr Gly Leu Ile Leu Thr  
 390 395 400  
 AGA GAT GGT GGT AAC AAC AAC ATG AGC AAG ACC ACC GAG 1245  
 Arg Asp Gly Gly Asn Asn Asn Met Ser Lys Thr Thr Glu  
 405 410 415  
 ACC TTC AGA CCT GGA GGA GGA GAT ATG AGG GAC AAT TGG 1284  
 Thr Phe Arg Pro Gly Gly Asp Met Arg Asp Asn Trp  
 420 425  
 AGA AGT GAA TTA TAT AAA TAT AAA GCA CTA AAA ATT GAA 1323  
 Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile Glu  
 430 435 440  
 CCA TTA GGA GCA GCA CCC ACC AGG GCA AAG AGA AGA GTG 1362  
 Pro Leu Gly Val Ala Pro Thr Arg Ala Lys Arg Arg Val  
 445 450  
 GTG CAG AGA GAA AAA AGA GCA GTG GGA ATA GGA GCT GTG 1401  
 Val Gln Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Val  
 455 460 465  
 TTC CTT GGG TTC TTG GGA GCA TAA AGC TTC TAG A 1435  
 Phe Leu Gly Phe Leu Gly Ala Xaa Ser Phe Xaa  
 470 475 478

## (2) INFORMATION FOR SEQ ID NO:26:

## (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 478 amino acids

(B) TYPE: Amino Acid

(D) TOPOLOGY: Linear

## 5 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:26:

Leu	Glu	Val	Pro	Val	Trp	Lys	Glu	Ala	Thr	Thr	Thr	Leu	Phe	Cys	
1															
														15	
10	Ala	Ser	Asp	Ala	Lys	Ala	Tyr	Asp	Ser	Glu	Ala	His	Asn	Val	Trp
														30	
15	Ala	Thr	His	Ala	Cys	Val	Pro	Thr	Asp	Pro	Asn	Pro	Gln	Glu	Val
														45	
20	Glu	Leu	Glu	Asn	Val	Thr	Glu	Asn	Phe	Asn	Met	Trp	Lys	Asn	Asn
														60	
25	Met	Val	Glu	Gln	Met	His	Gly	Asp	Ile	Ile	Ser	Leu	Trp	Asp	Gln
														75	
30	Ser	Leu	Lys	Pro	Cys	Val	Lys	Leu	Thr	Pro	Leu	Cys	Val	Thr	Leu
														90	
35	Asn	Cys	Thr	Asp	Pro	Asn	Val	Thr	Asn	Ser	Glu	Arg	Thr	Ile	Glu
														105	
40	Gly	Gly	Glu	Ile	Lys	Asn	Cys	Ser	Phe	Asn	Ile	Thr	Thr	Asn	Ile
														120	
45	Arg	Asp	Arg	Phe	Gln	Lys	Glu	Tyr	Ala	Leu	Phe	Tyr	Lys	Leu	Asp
														135	
50	Val	Ile	Pro	Leu	Gly	Asn	Asp	Asn	Thr	Ser	Tyr	Arg	Leu	Ile	Ser
														150	
55	Cys	Asn	Thr	Ser	Val	Ile	Thr	Gln	Ala	Cys	Pro	Lys	Val	Ser	Phe
														165	
60	Glu	Pro	Ile	Pro	Ile	His	Tyr	Cys	Ala	Pro	Ala	Gly	Phe	Ala	Ile
														180	
65	Leu	Lys	Cys	Lys	Asp	Lys	Phe	Asn	Gly	Thr	Gly	Pro	Cys	Thr	
														195	
70	Asn	Val	Ser	Thr	Val	Gln	Cys	Thr	His	Gly	Ile	Lys	Pro	Val	Val
														210	
75	Ser	Thr	Gln	Leu	Leu	Asn	Gly	Ser	Leu	Ala	Glu	Glu	Asp	Ile	
														225	
80	Val	Ile	Arg	Ser	Ala	Asn	Leu	Thr	Asp	Asn	Ala	Lys	Asn	Ile	Ile
														240	
85	Val	Gln	Leu	Asn	Glu	Ser	Val	Thr	Met	Asn	Cys	Thr	Arg	Pro	Asn
														255	
90	Asn	Asn	Thr	Met	Lys	Ser	Ile	His	Ile	Gly	Pro	Gly	Arg	Ala	Phe
														270	
95	Tyr	Ala	Thr	Gly	Asn	Ile	Ile	Gly	Asp	Ile	Arg	Gln	Ala	His	Cys
														285	

Asn Ile Ser Gly Thr Lys Trp Asn Asp Thr Leu Lys Lys Ile Ala  
 290 295 300  
 Ile Lys Leu Arg Glu Gln Phe Asn Lys Thr Ile Val Phe Asn Gln  
 5 305 310 315  
 Ser Ser Gly Gly Asp Pro Glu Ile Ala Thr Leu Ser Phe Asn Cys  
 320 325 330  
 10 Gly Gly Glu Phe Phe Tyr Cys Asn Ser Thr Gln Leu Phe Asn Ser  
 335 340 345  
 Thr Trp Asn Ser Thr Gly Ser Asn Asn Thr Lys Gly Asn Asp Thr  
 350 355 360  
 15 Ile Thr Leu Pro Cys Arg Ile Arg Gln Ile Ile Asn Met Trp Gln  
 365 370 375  
 Lys Ile Gly Lys Ala Met Tyr Ala Pro Pro Ile Lys Gly Gln Ile  
 20 380 385 390  
 Arg Cys Ser Ser Asn Ile Thr Gly Leu Ile Leu Thr Arg Asp Gly  
 395 400 405  
 25 Gly Asn Asn Asn Met Ser Lys Thr Thr Glu Thr Phe Arg Pro Gly  
 410 415 420  
 Gly Gly Asp Met Arg Asp Asn Trp Arg Ser Glu Leu Tyr Lys Tyr  
 425 430 435  
 30 Lys Val Val Lys Ile Glu Pro Leu Gly Val Ala Pro Thr Arg Ala  
 440 445 450  
 Lys Arg Arg Val Val Gln Arg Glu Lys Arg Ala Val Gly Ile Gly  
 35 455 460 465  
 Ala Val Phe Leu Gly Phe Leu Gly Ala Xaa Ser Phe Xaa  
 470 475 478

## 40 (2) INFORMATION FOR SEQ ID NO:27:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 1435 base pairs
- (B) TYPE: Nucleic Acid
- (C) STRANDEDNESS: Single

## 45 (D) TOPOLOGY: Linear

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:27:

CTC GAG GTA CCT GTG TGG AAA GAA GCA ACC ACC ACT 36  
 Leu Glu Val Pro Val Trp Lys Glu Ala Thr Thr Thr  
 50 1 5 10  
 CTA TTT TGT GCA TCA GAT GCT AAA GCA TAT GAT TCA GAG 75  
 Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Ser Glu  
 15 20 25  
 55 GCA CAT AAT GTT TGG GCC ACA CAT GCC TGT GTA CCC ACA 114  
 Ala His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr  
 30 35  
 60 GAC CCC AAC CCA CAA GAA GCA GAA TTG GAA AAT GTG ACA 153  
 Asp Pro Asn Pro Gln Glu Val Glu Leu Glu Asn Val Thr  
 40 45 50

GAA AAT TTT AAC ATG TGG AAA AAT AAC ATG GTA GAA CAG 192  
 Glu Asn Phe Asn Met Trp Lys Asn Asn Met Val Glu Gln  
 55 60

5 ATG CAT GGG GAT ATA ATT AGT TTA TCG GAT CAA AGC CTA 231  
 Met His Gly Asp Ile Ile Ser Leu Trp Asp Gln Ser Leu  
 65 70 75

10 AAG CCA TGT GTA AAA TTA ACC CCA CTC TGT GTT ACG TTA 270  
 Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu  
 80 85 90

15 AAT TGC ACT GAC CCA AAT GTT ACT AAT ACC GAG AGA ACG 309  
 Asn Cys Thr Asp Pro Asn Val Thr Asn Ser Glu Arg Thr  
 95 100

20 ATA GAG GGG GGA GAA ATA AAA AAT TGC TCT TTC AAT ATC 348  
 Ile Glu Gly Gly Glu Ile Lys Asn Cys Ser Phe Asn Ile  
 105 110 115

25 ACC ACA AAC ATA AGA GAT AGG TTT CAG AAA GAA TAT GCA 387  
 Thr Thr Asn Ile Arg Asp Arg Phe Gln Lys Glu Tyr Ala  
 120 125

30 CTT TTT TAT AAA CTT GAT GTA ATA CCA TTA CGT AAT GAT 426  
 Leu Phe Tyr Lys Leu Asp Val Ile Pro Leu Gly Asn Asp  
 130 135 140

35 AAT ACT AGC TAT AGG TTG ATA AGT TGT AAC ACC TCA GTC 465  
 Asn Thr Ser Tyr Arg Leu Ile Ser Cys Asn Thr Ser Val  
 145 150 155

40 ATT ACA CAG GCC TGT CCA AAG GTA TCC TTT GAG CCA ATT 504  
 Ile Thr Gln Ala Cys Pro Lys Val Ser Phe Glu Pro Ile  
 160 165

45 CCC ATA CAT TAT TGT GCC CCG GCT GGT TTT GCG ATT CTA 543  
 Pro Ile His Tyr Cys Ala Pro Ala Gly Phe Ala Ile Leu  
 170 175 180

50 AAG TGT AAA GAT AAG AAG TTC AAT GGA ACA GGA CCA TGT 582  
 Lys Cys Lys Asp Lys Lys Phe Asn Gly Thr Gly Pro Cys  
 185 190

55 ACA AAT GTC AGC ACA GTA CAA TGT ACA CAT GGA ATT AAG 621  
 Thr Asn Val Ser Thr Val Gln Cys Thr His Gly Ile Lys  
 195 200 205

60 CCA GTA GTA TCA ACT CAA CTG TTG TTA AAT GGC AGT CTA 660  
 Pro Val Val Ser Thr Gln Leu Leu Asn Gly Ser Leu  
 210 215 220

65 GCA GAA GAA GAC ATA GTA ATT AGA TCC GCC AAT CTC ACA 699  
 Ala Glu Glu Asp Ile Val Ile Arg Ser Ala Asn Leu Thr  
 225 230

70 GAC AAT GCT AAA AAC ATA ATA GTA CAG CTG AAT GAA TCT 738  
 Asp Asn Ala Lys Asn Ile Ile Val Gln Leu Asn Glu Ser  
 235 240 245

75 GTA ACA ATG AAT TGT ACA AGA CCC AAC AAC AAT ACA ATG 777  
 Val Thr Met Asn Cys Thr Arg Pro Asn Asn Asn Thr Met  
 250 255

AAA AGT ATA CAT ATA GGA CCA GGC AGA GCA TTT TAT GCA 816  
 Lys Ser Ile His Ile Gly Pro Gly Arg Ala Phe Tyr Ala  
 260 265 270  
 5 ACA GGA AAC ATA ATA GGA GAT ATA AGA CAA GCA CAT TGT 855  
 Thr Gly Asn Ile Ile Gly Asp Ile Arg Gln Ala His Cys  
 275 280 285  
 AAC ATT AGT GGA ACA AAA TGG AAT GAC ACT TTG AAA AAG 894  
 10 Asn Ile Ser Gly Thr Lys Trp Asn Asp Thr Leu Lys Lys  
 290 295  
 ATA GCT ATA AAA TTA AGA GAA CAA TTT AAT AAG ACA ATA 933  
 Ile Ala Ile Lys Leu Arg Glu Gln Phe Asn Lys Thr Ile  
 15 300 305 310  
 GTC TTT AAT CAA TCC TCA CGA GGG GAC CCA GAA ATT GCA 972  
 Val Phe Asn Gln Ser Ser Gly Gly Asp Pro Glu Ile Ala  
 315 320  
 20 ACG CTC AGT TTT AAT TGT GGA GGG GAA TTT TTC TAC TGT 1011  
 Thr Leu Ser Phe Asn Cys Gly Gly Glu Phe Phe Tyr Cys  
 325 330 335  
 25 AAT TCA ACA CAA CTG TTT AAT AGT ACT TGG AAT AGT ACT 1050  
 Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp Asn Ser Thr  
 340 345 350  
 30 GGG TCA AAT AAC ACT AAA GGA AAT GAC ACA ATC ACA CTC 1089  
 Gly Ser Asn Asn Thr Lys Gly Asn Asp Thr Ile Thr Leu  
 355 360  
 35 CCA TGC AGA ATA AGA CAA ATT ATA AAC ATG TGG CAG AAA 1128  
 Pro Cys Arg Ile Arg Gln Ile Ile Asn Met Trp Gln Lys  
 365 370 375  
 ATA GGA AAA GCA ATG TAT GCC CCT CCC ATC AAA GGG CAA 1167  
 Ile Gly Lys Ala Met Tyr Ala Pro Pro Ile Lys Gly Gln  
 380 385  
 40 ATT AGA TGT TCA TCA AAT ATT ACA CGG CTA ATA TTA ACA 1206  
 Ile Arg Cys Ser Ser Asn Ile Thr Gly Leu Ile Leu Thr  
 390 395 400  
 45 AGA CAT GGT GGT AAC AAC AAC ATG AGC AAG ACC ACC GAG 1245  
 Arg Asp Gly Gly Asn Asn Asn Met Ser Lys Thr Thr Glu  
 405 410 415  
 50 ACC TTC AGA CCT GGA GGA GGA GAT ATG AGG GAC AAT TGG 1284  
 Thr Phe Arg Pro Gly Gly Asp Met Arg Asp Asn Trp  
 420 425  
 55 AGA AGT GAA TTA TAT AAA TAT AAA GTA GTA AAA ATT GAA 1323  
 Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile Glu  
 430 435 440  
 CCA TTA GGA GTA GCA CCC ACC AGG GCA AAG AGA AGA GTG 1362  
 Pro Leu Gly Val Ala Pro Thr Arg Ala Lys Arg Arg Val  
 445 450  
 60 GTG CAG AGA GAA AAA AGA GCA GTG GGA ATA GGA GCT GTG 1401  
 Val Gln Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Val  
 455 460 465

TTC CTT GGG TTC TTG GGA GCA TAA AGC TTC TAG A 1435  
 Phe Leu Gly Phe Leu Gly Ala Xaa Ser Phe Xaa  
 470 475 478

## 5 (2) INFORMATION FOR SEQ ID NO:28:

(i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 478 amino acids  
 (B) TYPE: Amino Acid  
 (D) TOPOLOGY: Linear

## 10 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:28:

Leu Glu Val Pro Val Trp Lys Glu Ala Thr Thr Thr Leu Phe Cys  
 1 5 10 15

15 Ala Ser Asp Ala Lys Ala Tyr Asp Ser Glu Ala His Asn Val Trp  
 20 25 30

Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn Pro Gln Glu Val  
 35 40 45

20 Glu Leu Glu Asn Val Thr Glu Asn Phe Asn Met Trp Lys Asn Asn  
 50 55 60

25 Met Val Glu Gln Met His Gly Asp Ile Ile Ser Leu Trp Asp Gln  
 65 70 75

Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu  
 80 85 90

30 Asn Cys Thr Asp Pro Asn Val Thr Asn Ser Glu Arg Thr Ile Glu  
 95 100 105

Gly Gly Glu Ile Lys Asn Cys Ser Phe Asn Ile Thr Thr Asn Ile  
 110 115 120

35 Arg Asp Arg Phe Gln Lys Glu Tyr Ala Leu Phe Tyr Lys Leu Asp  
 125 130 135

40 Val Ile Pro Leu Gly Asn Asp Asn Thr Ser Tyr Arg Leu Ile Ser  
 140 145 150

Cys Asn Thr Ser Val Ile Thr Gln Ala Cys Pro Lys Val Ser Phe  
 155 160 165

45 Glu Pro Ile Pro Ile His Tyr Cys Ala Pro Ala Gly Phe Ala Ile  
 170 175 180

Leu Lys Cys Lys Asp Lys Lys Phe Asn Gly Thr Gly Pro Cys Thr  
 185 190 195

50 Asn Val Ser Thr Val Gln Cys Thr His Gly Ile Lys Pro Val Val  
 200 205 210

Ser Thr Gln Leu Leu Asn Gly Ser Leu Ala Glu Glu Asp Ile  
 215 220 225

Val Ile Arg Ser Ala Asn Leu Thr Asp Asn Ala Lys Asn Ile Ile  
 230 235 240

60 Val Gln Leu Asn Glu Ser Val Thr Met Asn Cys Thr Arg Pro Asn  
 245 250 255

Asn Asn Thr Met Lys Ser Ile His Ile Gly Pro Gly Arg Ala Phe  
 260 265 270

65

Tyr Ala Thr Gly Asn Ile Ile Gly Asp Ile Arg Gln Ala His Cys  
 275 280 285  
 Asn Ile Ser Gly Thr Lys Trp Asn Asp Thr Leu Lys Lys Ile Ala  
 5 290 295 300  
 Ile Lys Leu Arg Glu Gln Phe Asn Lys Thr Ile Val Phe Asn Gln  
 305 310 315  
 10 Ser Ser Gly Gly Asp Pro Glu Ile Ala Thr Leu Ser Phe Asn Cys  
 320 325 330  
 Gly Gly Glu Phe Phe Tyr Cys Asn Ser Thr Gln Leu Phe Asn Ser  
 335 340 345  
 15 Thr Trp Asn Ser Thr Gly Ser Asn Asn Thr Lys Gly Asn Asp Thr  
 350 355 360  
 Ile Thr Leu Pro Cys Arg Ile Arg Gln Ile Ile Asn Met Trp Gln  
 20 365 370 375  
 Lys Ile Gly Lys Ala Met Tyr Ala Pro Pro Ile Lys Gly Gln Ile  
 380 385 390  
 25 Arg Cys Ser Ser Asn Ile Thr Gly Leu Ile Leu Thr Arg Asp Gly  
 395 400 405  
 Gly Asn Asn Asn Met Ser Lys Thr Thr Glu Thr Phe Arg Pro Gly  
 410 415 420  
 30 Gly Gly Asp Met Arg Asp Asn Trp Arg Ser Glu Leu Tyr Lys Tyr  
 425 430 435  
 Lys Val Val Lys Ile Glu Pro Leu Gly Val Ala Pro Thr Arg Ala  
 35 440 445 450  
 Lys Arg Arg Val Val Gln Arg Glu Lys Arg Ala Val Gly Ile Gly  
 455 460 465  
 40 Ala Val Phe Leu Gly Phe Leu Gly Ala Xaa Ser Phe Xaa  
 470 475 478

## (2) INFORMATION FOR SEQ ID NO:29:

## (i) SEQUENCE CHARACTERISTICS:

45 (A) LENGTH: 511 amino acids  
 (B) TYPE: Amino Acid  
 (D) TOPOLOGY: Linear

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:29:

50 Met Arg Val Lys Gly Ile Arg Arg Asn Tyr Gln His Trp Trp Gly Arg  
 1 5 10 15  
 Gly Thr Met Leu Leu Gly Leu Leu Met Ile Cys Ser Ala Thr Glu Lys  
 20 25 30  
 55 Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala Thr  
 35 40 45  
 Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu Ala  
 60 50 55 60  
 His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn Pro  
 65 70 75 80

Gln Glu Val Glu Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp Lys  
 85 90 95  
 Asn Asn Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp Asn  
 5 100 105 110  
 Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu  
 115 120 125  
 10 Asn Cys Thr Asp Leu Arg Asn Thr Thr Asn Thr Asn Asn Ser Thr Asp  
 130 135 140  
 Asn Asn Asn Ser Lys Ser Glu Gly Thr Ile Lys Gly Gly Glu Met Lys  
 145 150 155 160  
 15 Asn Cys Ser Phe Asn Ile Thr Thr Ser Ile Gly Asp Lys Met Gln Lys  
 165 170 175  
 20 Glu Tyr Ala Leu Leu Tyr Lys Leu Asp Ile Glu Pro Ile Asp Asn Asp  
 180 185 190  
 Ser Thr Ser Tyr Arg Leu Ile Ser Cys Asn Thr Ser Val Ile Thr Gln  
 195 200 205  
 25 Ala Cys Pro Lys Ile Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala  
 210 215 220  
 Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asp Lys Lys Phe Ser Gly  
 225 230 235 240  
 30 Lys Gly Ser Cys Lys Asn Val Ser Thr Val Gln Cys Thr His Gly Ile  
 245 250 255  
 Arg Pro Val Val Ser Thr Gln Leu Leu Asn Gly Ser Leu Ala Glu  
 35 260 265 270  
 Glu Glu Val Val Ile Arg Ser Glu Asp Phe Thr Asp Asn Ala Lys Thr  
 275 280 285  
 40 Ile Ile Val His Leu Lys Glu Ser Val Gln Ile Asn Cys Thr Arg Pro  
 290 295 300  
 Asn Tyr Asn Lys Arg Lys Arg Ile His Ile Gly Pro Gly Arg Ala Phe  
 305 310 315 320  
 45 Tyr Thr Thr Lys Asn Ile Lys Gly Thr Ile Arg Gln Ala His Cys Ile  
 325 330 335  
 Ile Ser Arg Ala Lys Trp Asn Asp Thr Leu Arg Gln Ile Val Ser Lys  
 50 340 345 350  
 Leu Lys Glu Gln Phe Lys Asn Lys Thr Ile Val Phe Asn Pro Ser Ser  
 355 360 365  
 55 Gly Gly Asp Pro Glu Ile Val Met His Ser Phe Asn Cys Gly Gly Glu  
 370 375 380  
 Phe Phe Tyr Cys Asn Thr Ser Pro Leu Phe Asn Ser Ile Trp Asn Gly  
 385 390 395 400  
 60 Asn Asn Thr Trp Asn Asn Thr Thr Gly Ser Asn Asn Asn Ile Thr Leu  
 405 410 415  
 Gln Cys Lys Ile Lys Gln Ile Ile Asn Met Trp Gln Lys Val Gly Lys  
 65 420 425 430

	Ala	Met	Tyr	Ala	Pro	Pro	Ile	Glu	Gly	Gln	Ile	Arg	Cys	Ser	Ser	Asn
435							440					445				
	Ile	Thr	Gly	Leu	Leu	Leu	Thr	Arg	Asp	Gly	Gly	Glu	Asp	Thr	Asp	Thr
5	450						455					460				
	Asn	Asp	Thr	Glu	Ile	Phe	Arg	Pro	Gly	Gly	Gly	Asp	Met	Arg	Asp	Asn
465						470					475			480		
10	Trp	Arg	Ser	Glu	Leu	Tyr	Lys	Tyr	Lys	Val	Val	Thr	Ile	Glu	Pro	Leu
							485			490			495			
	Gly	Val	Ala	Pro	Thr	Lys	Ala	Lys	Arg	Arg	Val	Val	Gln	Arg	Glu	
15							500			505			510			

## (2) INFORMATION FOR SEQ ID NO:30

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 2800 base pairs
- (B) TYPE: Nucleic Acid
- (C) STRANDEDNESS: Single
- (D) TOPOLOGY: Linear

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:30:

25	TTCGAGCTCG	CCCGACATTG	ATTATTGACT	AGAGTCGATC	GACAGCTGTG	50									
	GAATGTGTGT	CAGTAGGGT	GTGGAAAGTC	CCCAAGGCTCC	CCAGCAGGCA	100									
30	GAACATATGCA	AAGCATGCAT	CTCAATTAGT	CAGCAACCAG	GTGTGGAAAG	150									
	TCCCCAGGCT	CCCCAGCAGG	CAGAACTATG	CAAAGCATGC	ATCTCAATTA	200									
	GTCAGCAACC	ATAGTCCCAG	CCCTAACTCC	GCCCATCCCG	CCCCTAACTC	250									
35	CGCCCAGTTC	CGCCCATTC	CCGCCCCATG	GCTGACTAAT	TTTTTTTATT	300									
	TATGCAGAGG	CCGAGGCCGC	CTCGGCCTCT	GAGCTATTCC	AGAACTAGTG	350									
	AGGAGGCCTT	TTTGGAGGCC	TAGGCTTTG	AAAAAGCTA	GCTTATCCCG	400									
40	CCGGGAACGG	TGCATTGGAA	CCGGGATTCC	CCGTGCCAAG	AGTCAGGTAA	450									
	GTACCGCCTA	TAGAGTCTAT	AGGCCCACCC	CCTTGGCTTC	GTAGAACCGC	500									
45	GGCTACAATT	AATACATAAC	CTTTGGATC	GATCCTACTG	ACACTGACAT	550									
	CCACTTTTC	TTTTCTCCA	CAGGTGTCCA	CTCCCAGGTC	CAACTGCACC	600									
	TCGGTTCGCG	AAGCTAGCTT	GGGCTGCATC	GATTGAATTG	CACTGCCTTC	650									
50	CACCAAGCTC	TGCAGGATCC	CAGAGTCAGG	GG	TCT GTC TCT TCC TGC	697									
				Ser Val Ser Ser Cys											
			1		5										
55	TGG	TGG	CTC	CAG	TTC	AGG	AAC	AGT	AAA	CCC	TGC	TCC	GAA	TAT	739
	Trp	Trp	Leu	Gln	Phe	Arg	Asn	Ser	Lys	Pro	Cys	Ser	Glu	Tyr	
	10													15	
60	TGC	CTC	TCA	CAT	CTC	GTC	AAT	CTC	CGC	GAG	GAC	TGG	GGG	CCC	781
	Cys	Leu	Ser	His	Leu	Val	Asn	Leu	Arg	Glu	Asp	Trp	Gly	Pro	
	20													30	
	25														

35	TCT GAC AAG CTT CAG CGC GAA CGA CCA ACT ACC CCG ATC ATC	823	
	Cys Asp Lys Leu Gln Arg Glu Arg Pro Thr Thr Pro Ile Ile		
40	45		
5	AGT TAT CCT TAA GGT CTC TTT TGT GTG CGT TCC GGT ATG	865	
Ser Tyr Pro	* Gly Leu Phe Cys Val Val Arg Ser Gly Met		
50	55	60	
10	GGG GGG ACT GCC GCC AGG TTG GGG GCC GTG ATT TTG TTT GTC	907	
Gly Gly Thr Ala Ala Arg Leu Gly Ala Val Ile Leu Phe Val			
62	65	70	75
15	GTC ATA GTG GGC CTC CAT GGG GTC CCC GGC AAA TAT GCC TTG	949	
Val Ile Val Gly Leu His Gly Val Arg Gly Lys Tyr Ala Leu			
20	80	85	
25	GCG GAT GCC TCT CTC AAG ATG GCC GAC CCC AAT CGA TTT CGC	991	
Ala Asp Ala Ser Leu Lys Met Ala Asp Pro Asn Arg Phe Arg			
90	95	100	
30	GGC AAA GAC CTT CCG GTC CTG GAC CAG CTG CTC GAG GTC CCT	1033	
Gly Lys Asp Leu Pro Val Leu Asp Gln Leu Leu Glu Val Pro			
105	110	115	
35	GTC TGG AAA GAA GCA AAC ACC ACT CTA TTT TGT GCA TCA GAT	1075	
Val Trp Lys Glu Ala Asn Thr Thr Leu Phe Cys Ala Ser Asp			
120	125	130	
40	GCT AAA GCA TAT AAG ACA GAG GCA CAT AAT GTT TCG GCC ACA	1117	
Ala Lys Ala Tyr Lys Thr Glu Ala His Asn Val Trp Ala Thr			
135	140	145	
45	CAT GCC TGT GTA CCC ACA GAC CCC AAA CCA CAA GAA ATA AAA	1159	
His Ala Cys Val Pro Thr Asp Pro Lys Pro Gln Glu Ile Lys			
150	155		
50	TTG GAA AAT GTG ACA GAA AAT TTT AAC ATG TGG AAA AAT AAC	1201	
Leu Glu Asn Val Thr Glu Asn Phe Asn Met Trp Lys Asn Asn			
160	165	170	
55	ATG GTA GAA CAG ATG CAT GAG GAT ATA ATC ACT TTA TGG GAT	1243	
Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp Asp			
175	180	185	
60	CAA AGC CTA AAG CCA TGT GTA AAA TTA ACC CCA CTC TGT GTT	1285	
Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val			
190	195	200	
65	ACT TTA AAT TGC ACT GAT TTG AGG AAT AAT ACT AAT ACC AAT	1327	
Thr Leu Asn Cys Thr Asp Leu Arg Asn Asn Thr Asn Thr Asn			
205	210	215	
70	AGT ACC TAC GGA AAA ATA ATG GAG GGA GGA GAG ATA AAA AAC	1369	
Ser Thr Tyr Gly Lys Ile Met Glu Gly Gly Glu Ile Lys Asn			
220	225		
75	TGC TCT TTC AAT ATC ACC ACA AGC ATA AAA GAT AAG CTC AAA	1411	
Cys Ser Phe Asn Ile Thr Thr Ser Ile Lys Asp Lys Leu Lys			
230	235	240	
80	GAT ATG TCA CTT TTT TAT AAA CTT GAT GTA GTA CCA ATA GGT	1453	
Asp Met Ser Leu Phe Tyr Lys Leu Asp Val Val Pro Ile Gly			
245	250	255	

AAT	AAT	AGT	AAT	ACT	ACT	AGT	TAT	AGG	TTG	ATA	AGT	TGT	AAC	1495	
Asn	Asn	Ser	Asn	Thr	Thr	Ser	Tyr	Arg	Leu	Ile	Ser	Cys	Asn		
260						265						270			
5	ACC	TCA	GTC	ATT	ACA	CAA	GCC	TGT	CCA	AAG	ACA	TCC	TTT	GAG	1537
	Thr	Ser	Val	Ile	Thr	Gln	Ala	Cys	Pro	Lys	Thr	Ser	Phe	Glu	
						275			280			285			
10	CCA	ATT	CCC	ATA	CAT	TAT	TGT	GCC	CCG	GCT	GGT	TTT	GCG	ATT	1579
	Pro	Ile	Pro	Ile	His	Tyr	Cys	Ala	Pro	Ala	Gly	Phe	Ala	Ile	
						290			295						
15	CTC	AAG	TGT	AAT	GAT	AAT	AAG	TTC	AAT	GGA	ACA	GGA	CCA	TGT	1621
	Leu	Lys	Cys	Asn	Asp	Asn	Lys	Phe	Asn	Gly	Thr	Gly	Pro	Cys	
						300		305		310					
20	CCA	AAT	GTC	AGC	ACA	GTA	CAA	TGT	ACA	CAT	GGA	ATT	AGG	CCA	1663
	Pro	Asn	Val	Ser	Thr	Val	Gln	Cys	Thr	His	Gly	Ile	Arg	Pro	
						315		320		325					
	GTA	GTA	TCA	ACT	CAA	CTG	CTG	TTA	AAT	GCC	AGT	CTA	GCA	GAA	1705
	Val	Val	Ser	Thr	Gln	Leu	Leu	Leu	Asn	Gly	Ser	Leu	Ala	Glu	
						330		335		340					
25	AAA	GAG	GTA	GTC	CTT	AGA	TCT	GAA	AAT	TTC	ACG	GAC	AAT	GCT	1747
	Lys	Glu	Val	Val	Val	Leu	Arg	Ser	Glu	Asn	Phe	Thr	Asp	Asn	
						345		350		355					
30	AAA	ACC	ATA	ATA	GTA	CAG	CTG	AAC	GAA	TCT	GTA	ATA	ATT	GAT	1789
	Lys	Thr	Ile	Ile	Val	Gln	Leu	Asn	Glu	Ser	Val	Ile	Ile	Asp	
						360		365							
35	TGT	ATG	AGA	CCC	AAC	AAC	AAT	ACA	AGA	ACA	AGT	ATA	CCT	ATG	1831
	Cys	Met	Arg	Pro	Asn	Asn	Asn	Thr	Arg	Thr	Ser	Ile	Pro	Met	
						370		375		380					
40	GGA	CCA	GGG	AAA	GCA	TTT	TAT	GCA	ACA	GGA	GAT	GTA	ATA	GGA	1873
	Gly	Pro	Gly	Lys	Ala	Phe	Tyr	Ala	Thr	Gly	Asp	Val	Ile	Gly	
						385		390		395					
	GAT	ATA	AGA	CGA	GCA	CAT	TGT	AAC	ATT	AGT	AGA	GCA	GGA	TGG	1915
	Asp	Ile	Arg	Arg	Ala	His	Cys	Asn	Ile	Ser	Arg	Ala	Gly	Trp	
						400		405		410					
45	AAT	ACC	ACT	TTA	CAA	CAG	ATA	GCT	AAA	AAA	TTA	AGA	GAA	AAA	1957
	Asn	Thr	Thr	Leu	Gln	Gln	Ile	Ala	Lys	Lys	Leu	Arg	Glu	Lys	
						415		420		425					
50	TTT	GAG	AAC	AAA	ACA	ATA	GTT	TTT	AAT	CAC	TCC	TCA	GGA	GGG	1999
	Phe	Glu	Asn	Lys	Thr	Ile	Val	Phe	Asn	His	Ser	Ser	Gly	Gly	
						430		435							
55	GAC	CCA	GAA	ATT	GTA	ATG	CAC	ACT	TTT	AAT	TGT	GGA	GGG	GAA	2041
	Asp	Pro	Glu	Ile	Val	Met	His	Thr	Phe	Asn	Cys	Gly	Gly	Glu	
						440		445		450					
60	TTT	TTC	TGC	TGT	AAT	TCA	ACA	CCA	CTG	TTT	AAT	AGT	ACT	TGG	2083
	Phe	Phe	Cys	Cys	Asn	Ser	Thr	Pro	Leu	Phe	Asn	Ser	Thr	Trp	
						455		460		465					
	AAT	GAT	GCA	CAA	CTG	TTT	AAT	AGT	ACT	TGG	GAT	GAT	ACT	AAA	2125
	Asn	Asp	Ala	Gln	Leu	Phe	Asn	Ser	Thr	Trp	Asp	Asp	Thr	Lys	
						470		475		480					
65															

10	TCG TCA AAA GGC ACT AAC GAA AAT GAC ACA ATC ACC CTC CAT Trp Ser Lys Gly Thr Asn Glu Asn Asp Thr Ile Thr Leu His 485 490 495	2167
5	TGG AGA ATA AAA CAA ATT ATA AAT ATG TGG CAG GAA GTA GGA Cys Arg Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly 500 505	2209
10	AAA GCA ATG TAT GCC CCT CCC ATC AAA GGA CAA ATT AGA TGT Lys Ala Met Tyr Ala Pro Pro Ile Lys Gly Gln Ile Arg Cys 510 515 520	2251
15	GAA TCA AAT ATT ACA GGG CTG CTA TTA ACA AGA GAT GGT GGT Glu Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly 525 530 535	2293
20	AAC GAC ACG AGC AAG AAT AAC ACT GAG ATT TTC AGA CCT GGA Asn Asp Thr Ser Lys Asn Asn Thr Glu Ile Phe Arg Pro Gly 540 545 550	2335
25	GGA GGA AAT ATG AAG GAC AAT TCG AGA ACT GAA TTA TAT AAA Gly Gly Asn Met Lys Asp Asn Trp Arg Ser Glu Leu Tyr Lys 555 560 565	2377
30	TAT AAA GTA ATA AAA ATT GAA CCA TTA GGA GTA GCA CCC ATC Tyr Lys Val Ile Lys Ile Glu Pro Leu Gly Val Ala Pro Ile 570 575 579	2419
35	TAGGCAAAGA GAAGAGTGGT GCAGAGAGAA AAAAGAGCAG TGACACTACG AGCTATGTTTC CTTGGGTTCT TGGGAGCAGC AGGAAGCACT ATGGGCATA AGCTTTAATG CGGTAGTTA TCACAGTTAA ATTCTGTAACG CACTCAGGCA CCGTGTATGA AATCTAACAA TGGCACCTCC AGAAGCTTAG AACCGAGGAA 40	2469 2519 2569 2619
45	CTTGTATTATT GCAGCTTATA ATGTTACAA ATAAACCAAT ACCATCACAA ATTCACAAA TAAAGCATT TTTCACTGC ATTCTAGTTG TCGTTGTCC AAACTCATCA ATGTATCTTA TCATGTCTGG ATCGGGATT AATTGGCGC AGCACCATGG CCTGAAATAA CCTCTGAAAG A	2669 2719 2769 2800

## 50 (2) INFORMATION FOR SEQ ID NO:31

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 579 amino acids
- (B) TYPE: Amino Acid
- (D) TOPOLOGY: Linear

## 55 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:31:

Ser Val Ser Ser Cys Trp Trp Leu Gln Phe Arg Asn Ser Lys Pro Cys	1 5 10 15
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60 Ser Glu Tyr Cys Leu Ser His Leu Val Asn Leu Arg Glu Asp Trp Gly	20 25 30
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Pro Cys Asp Lys Leu Gln Arg Glu Arg Pro Thr Thr Pro Ile Ile Ser	35 40 45
---	----------

Tyr Pro \* Gly Leu Phe Cys Val Val Arg Ser Gly Met Gly Gly Thr  
 50 55 60

Ala Ala Arg Leu Gly Ala Val Ile Leu Phe Val Val Ile Val Gly Leu  
 5 65 70 75 80

His Gly Val Arg Gly Lys Tyr Ala Leu Ala Asp Ala Ser Leu Lys Met  
 85 90 95

10 Ala Asp Pro Asn Arg Phe Arg Gly Lys Asp Leu Pro Val Leu Asp Gln  
 100 105 110

Leu Leu Glu Val Pro Val Trp Lys Glu Ala Asn Thr Thr Leu Phe Cys  
 115 120 125

15 Ala Ser Asp Ala Lys Ala Tyr Lys Thr Glu Ala His Asn Val Trp Ala  
 130 135 140

Thr His Ala Cys Val Pro Thr Asp Pro Lys Pro Gln Glu Ile Lys Leu  
 20 145 150 155 160

Glu Asn Val Thr Glu Asn Phe Asn Met Trp Lys Asn Asn Met Val Glu  
 165 170 175

25 Gln Met His Glu Asp Ile Ile Ser Leu Trp Asp Gln Ser Leu Lys Pro  
 180 185 190

Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu Asn Cys Thr Asp Leu  
 195 200 205

30 Arg Asn Asn Thr Asn Thr Asn Ser Thr Tyr Gly Lys Ile Met Glu Gly  
 210 215 220

Gly Glu Ile Lys Asn Cys Ser Phe Asn Ile Thr Thr Ser Ile Lys Asp  
 35 225 230 235 240

Lys Leu Lys Asp Met Ser Leu Phe Tyr Lys Leu Asp Val Val Pro Ile  
 245 250 255

40 Gly Asn Asn Ser Asn Thr Thr Ser Tyr Arg Leu Ile Ser Cys Asn Thr  
 260 265 270

Ser Val Ile Thr Gln Ala Cys Pro Lys Thr Ser Phe Glu Pro Ile Pro  
 275 280 285

45 Ile His Tyr Cys Ala Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asp  
 290 295 300

Asn Lys Phe Asn Gly Thr Gly Pro Cys Pro Asn Val Ser Thr Val Gln  
 50 305 310 315 320

Cys Thr His Gly Ile Arg Pro Val Val Ser Thr Cln Leu Leu Leu Asn  
 325 330 335

55 Gly Ser Leu Ala Glu Lys Glu Val Val Leu Arg Ser Glu Asn Phe Thr  
 340 345 350

Asp Asn Ala Lys Thr Ile Ile Val Gln Leu Asn Glu Ser Val Ile Ile  
 355 360 365

60 Asp Cys Met Arg Pro Asn Asn Asn Thr Arg Thr Ser Ile Pro Met Gly  
 370 375 380

Pro Gly Lys Ala Phe Tyr Ala Thr Gly Asp Val Ile Gly Asp Ile Arg  
 65 385 390 395 400

Arg Ala His Cys Asn Ile Ser Arg Ala Gly Trp Asn Thr Thr Leu Gln  
 405 410 415  
 Gln Ile Ala Lys Lys Leu Arg Glu Lys Phe Glu Asn Lys Thr Ile Val  
 5 420 425 430  
 Phe Asn His Ser Ser Gly Gly Asp Pro Glu Ile Val Met His Thr Phe  
 435 440 445  
 10 Asn Cys Gly Gly Glu Phe Phe Cys Asn Ser Thr Pro Leu Phe Asn  
 450 455 460  
 Ser Thr Trp Asn Asp Ala Gln Leu Phe Asn Ser Thr Trp Asp Asp Thr  
 465 470 475 480  
 15 Lys Trp Ser Lys Gly Thr Asn Glu Asn Asp Thr Ile Thr Leu His Cys  
 485 490 495  
 Arg Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys Ala Met  
 20 500 505 510  
 Tyr Ala Pro Pro Ile Lys Gly Gln Ile Arg Cys Glu Ser Asn Ile Thr  
 515 520 525  
 25 Gly Leu Leu Leu Thr Arg Asp Gly Gly Asn Asp Thr Ser Lys Asn Asn  
 530 535 540  
 Thr Glu Ile Phe Arg Pro Gly Gly Asn Met Lys Asp Asn Trp Arg  
 545 550 555 560  
 30 Ser Glu Leu Tyr Lys Tyr Lys Val Ile Lys Ile Glu Pro Leu Gly Val  
 565 570 575  
 Ala Pro Ile  
 35

## (2) INFORMATION FOR SEQ ID NO:32:

## (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1533 base pairs

(B) TYPE: Nucleic Acid

(C) STRANDEDNESS: Single

(D) TOPOLOGY: Linear

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:32:

45 ATGGGGGGGA CTGCCGCCAG GTTGGGGGCC GTGATTTGT TTGTCGTCAT 50  
 AGTGGGCCTC CATGGGGTCC GCGGCAAATA TGCCTTGGCG GATGCCCTCTC 100  
 TCAAGATGGC CGACCCCCAT CGATTTCGCG GCAAAGACCT TCCGGTCCTG 150  
 50 GACCAGCTGC TCGAG GTA CCT GTG TGG AAA GAA GCA ACC ACC 192  
 Val Pro Val Trp Lys Glu Ala Thr Thr  
 1 5  
 55 ACT CTA TTT TGT GCA TCA GAT GCT AAA GCA TAT GAT ACA GAG 234  
 Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu  
 10 15 20  
 60 GTA CAT AAT GTT TGG GCC ACA CAT GCC TGT GTA CCC ACA GAC 276  
 Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp  
 25 30 35

CCC AAC CCA CAA GAA ATA GGA TTG GAA AAT GTC ACA GAA AAT 318  
 Pro Asn Pro Gln Glu Ile Gly Leu Glu Asn Val Thr Glu Asn  
 40 45 50  
 5 TTT AAC ATC TCG AAA AAT AAC ATG GTC GAA CAG ATG CAT GAG 360  
 Phe Asn Met Trp Lys Asn Asn Met Val Glu Gln Met His Glu  
 55 60 65  
 GAT ATA ATC AGT TTA TGG GAT CAA AGC TTA AAG CCA TGT GTC 402  
 10 Asp Ile Ile Ser Leu Trp Asp Gln Ser Leu Lys Pro Cys Val  
 70 75  
 AAA TTA ACC CCA CTA TGT GTT ACT TTA AAT TGC ACT GAT TTG 444  
 Lys Leu Thr Pro Leu Cys Val Thr Leu Asn Cys Thr Asp Leu  
 15 80 85 90  
 20 AAA AAT GCT ACT AAT ACC ACT AGT AGC AGC TGG GGA AAG ATG 486  
 Lys Asn Ala Thr Asn Thr Ser Ser Ser Trp Gly Lys Met  
 95 100 105  
 GAG AGA GGA GAA ATA AAA AAC TGC TCT TTC AAT GTC ACC ACA 528  
 Glu Arg Gly Glu Ile Lys Asn Cys Ser Phe Asn Val Thr Thr  
 110 115 120  
 25 ACT ATA AGA GAT AAG ATG AAG AAT GAA TAT GCA CTT TTT TAT 570  
 Ser Ile Arg Asp Lys Met Lys Asn Glu Tyr Ala Leu Phe Tyr  
 125 130 135  
 30 AAA CTT GAT GTA GTC CCA ATA GAT AAT GAT AAT ACT AGC TAT 612  
 Lys Leu Asp Val Val Pro Ile Asp Asn Asp Asn Thr Ser Tyr  
 140 145  
 AGG TTG ATA AGT TGT AAC ACC TCA GTC ATT ACA CAG GCC TGT 654  
 35 Arg Leu Ile Ser Cys Asn Thr Ser Val Ile Thr Glu Ala Cys  
 150 155 160  
 CCA AAG GTG TCC TTT GAG CCA ATT CCC ATA CAT TAT TGT GCC 696  
 40 Pro Lys Val Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala  
 165 170 175  
 CCG GCT GGT TTT GCG ATT CTA AAG TGT AGA GAT AAA AAG TTC 738  
 Pro Ala Gly Phe Ala Ile Leu Lys Cys Arg Asp Lys Lys Phe  
 45 180 185 190  
 AAC GGA ACA GGA CCA TGT ACA AAT GTC AGC ACA GTC CAA TGT 780  
 Asn Gly Thr Gly Pro Cys Thr Asn Val Ser Thr Val Glu Cys  
 195 200 205  
 50 ACA CAT GGA ATT AGG CCA GTC GTC TCA ACT CAA CTG CTG TTA 822  
 Thr His Gly Ile Arg Pro Val Val Ser Thr Glu Leu Leu  
 210 215  
 55 AAT GGC AGT TTA GCA GAA GAA GAA GTC GTC ATT ACA TCT GCC 864  
 Asn Gly Ser Leu Ala Glu Glu Val Val Ile Arg Ser Ala  
 220 225 230  
 60 AAT TTC TCG GAC AAT GCT AAA ACC ATA ATA GTC CAG CTG AAC 906  
 Asn Phe Ser Asp Asn Ala Lys Thr Ile Ile Val Glu Leu Asn  
 235 240 245

10	GAA TCT GTA GAA ATT AAT TGT ACA AGA CCC AAC AAC AAT ACA Glu Ser Val Glu Ile Asn Cys Thr Arg Pro Asn Asn Asn Thr 250 255 260	948
5	AGA AGA AGT ATA CAT ATA GGA CCA GGG AGA GCA TTT TAT GCA Arg Arg Ser Ile His Ile Gly Pro Gly Arg Ala Phe Tyr Ala 265 270 275	990
10	ACA GGA GAA ATA ATA GGA GAC ATA AGA CAA GCA CAT TGT AAC Thr Gly Glu Ile Ile Gly Asp Ile Arg Gln Ala His Cys Asn 280 285	1032
15	CTT AGT AGC ACA AAA TGG AAT AAT ACT TTA AAA CAG ATA GTT Leu Ser Ser Thr Lys Trp Asn Asn Thr Leu Lys Gln Ile Val 290 295 300	1074
20	ACA AAA TTA AGA GAA CAT TTT AAT AAA ACA ATA GTC TTT AAT Thr Lys Leu Arg Glu His Phe Asn Lys Thr Ile Val Phe Asn 305 310 315	1116
25	CAC TCC TCA GGA GGG GAC CCA GAA ATT GTA ATG CAC AGT TTT His Ser Ser Gly Gly Asp Pro Glu Ile Val Met His Ser Phe 320 325 330	1158
30	AAT TGT GGA GGG GAA TTT TTC TAC TGT AAT ACA ACA CCA CTG Asn Cys Gly Glu Phe Phe Tyr Cys Asn Thr Thr Pro Leu 335 340 345	1200
35	TTT AAT AGT ACT TGG AAT TAT ACT TAT ACT TGG AAT AAT ACT Phe Asn Ser Thr Trp Asn Tyr Thr Tyr Thr Trp Asn Asn Thr 350 355	1242
40	GAA GGG TCA AAT GAC ACT GGA AGA AAT ATC ACA CTC CAA TGC Glu Gly Ser Asn Asp Thr Gly Arg Asn Ile Thr Leu Gln Cys 360 365 370	1284
45	AGA ATA AAA CAA ATT ATA AAC ATG TGG CAG GAA GCA GGA AAA Arg Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys 375 380 385	1326
50	GCA ATG TAT GCC CCT CCC ATA AGA GGA CAA ATT AGA TGC TCA Ala Met Tyr Ala Pro Pro Ile Arg Gly Gln Ile Arg Cys Ser 390 395 400	1368
55	TCA AAT ATT ACA GGG CTG CTA TTA ACA AGA GAT GGT GGT AAT Ser Asn Ile Thr Gly Leu Leu Thr Arg Asp Gly Gly Asn 405 410 415	1410
60	AAC AGC GAA ACC GAG ATC TTC AGA CCT GGA GGA GCA GAT ATG Asn Ser Glu Thr Glu Ile Phe Arg Pro Gly Gly Gly Asp Met 420 425	1452
65	AGG GAC AAT TGG AGA AGT GAA TTA TAT AAA TAT AAA GCA GCA Arg Asp Asn Trp Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val 430 435 440	1494
70	AAA ATT GAA CCA TTA GGA GTA GCA CCC ACC AAG GCA TAA Lys Ile Glu Pro Leu Gly Val Ala Pro Thr Lys Ala *	1533
75	445 450 455	

## (2) INFORMATION FOR SEQ ID NO:33:

## (1) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 456 amino acids

(B) TYPE: Amino Acid

(D) TOPOLOGY: Linear

## 5 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:33:

Val	Pro	Val	Trp	Lys	Glu	Ala	Thr	Thr	Thr	Leu	Phe	Cys	Ala	Ser	Asp		
1	5				10					15							
10		Ala	Lys	Ala	Tyr	Asp	Thr	Glu	Val	His	Asn	Val	Trp	Ala	Thr	His	Ala
		20				25					30						
15		Cys	Val	Pro	Thr	Asp	Pro	Asn	Pro	Gln	Glu	Ile	Gly	Leu	Glu	Asn	Val
		35				40					45						
20		Thr	Glu	Asn	Phe	Asn	Met	Trp	Lys	Asn	Asn	Met	Val	Glu	Gln	Met	His
		50				55					60						
25		Glu	Asp	Ile	Ile	Ser	Leu	Trp	Asp	Gln	Ser	Leu	Lys	Pro	Cys	Val	Lys
		65				70					75					80	
30		Leu	Thr	Pro	Leu	Cys	Val	Thr	Leu	Asn	Cys	Thr	Asp	Leu	Lys	Asn	Ala
		85								90						95	
35		Thr	Asn	Thr	Thr	Ser	Ser	Ser	Trp	Gly	Lys	Met	Glu	Arg	Gly	Glu	Ile
		100								105					110		
40		Lys	Asn	Cys	Ser	Phe	Asn	Val	Thr	Thr	Ser	Ile	Arg	Asp	Lys	Met	Lys
		115								120					125		
45		Asn	Glu	Tyr	Ala	Leu	Phe	Tyr	Lys	Leu	Asp	Val	Val	Pro	Ile	Asp	Asn
		130							135					140			
50		Asp	Asn	Thr	Ser	Tyr	Arg	Leu	Ile	Ser	Cys	Asn	Thr	Ser	Val	Ile	Thr
		145					150					155					160
55		Gln	Ala	Cys	Pro	Lys	Val	Ser	Phe	Glu	Pro	Ile	Pro	Ile	His	Tyr	Cys
		165							170						175		
60		Ala	Pro	Ala	Gly	Phe	Ala	Ile	Leu	Lys	Cys	Arg	Asp	Lys	Lys	Phe	Asn
		180							185						190		
65		Gly	Thr	Gly	Pro	Cys	Thr	Asn	Val	Ser	Thr	Val	Gln	Cys	Thr	His	Gly
		195							200						205		
70		Ile	Arg	Pro	Val	Val	Ser	Thr	Gln	Leu	Leu	Leu	Asn	Gly	Ser	Leu	Ala
		210								215					220		
75		Glu	Glu	Glu	Val	Val	Ile	Arg	Ser	Ala	Asn	Phe	Ser	Asp	Asn	Ala	Lys
		225							230						235		240
80		Thr	Ile	Ile	Val	Gln	Leu	Asn	Glu	Ser	Val	Glu	Ile	Asn	Cys	Thr	Arg
		245										250				255	
85		Pro	Asn	Asn	Asn	Thr	Arg	Arg	Ser	Ile	His	Ile	Gly	Pro	Gly	Arg	Ala
		260								265						270	
90		Phe	Tyr	Ala	Thr	Gly	Glu	Ile	Gly	Asp	Ile	Arg	Gln	Ala	His	Cys	
		275							280						285		
95		Asn	Leu	Ser	Ser	Thr	Lys	Trp	Asn	Asn	Thr	Leu	Lys	Gln	Ile	Val	Thr
		290							295						300		

Lys Leu Arg Glu His Phe Asn Lys Thr Ile Val Phe Asn His Ser Ser  
 305 310 315 320  
 Gly Gly Asp Pro Glu Ile Val Met His Ser Phe Asn Cys Gly Gly Glu  
 5 325 330 335  
 Phe Phe Tyr Cys Asn Thr Thr Pro Leu Phe Asn Ser Thr Trp Asn Tyr  
 340 345 350  
 10 Thr Tyr Thr Trp Asn Asn Thr Glu Gly Ser Asn Asp Thr Gly Arg Asn  
 355 360 365  
 Ile Thr Leu Gln Cys Arg Ile Lys Gln Ile Ile Asn Met Trp Gln Glu  
 15 370 375 380  
 Val Gly Lys Ala Met Tyr Ala Pro Pro Ile Arg Gly Gln Ile Arg Cys  
 385 390 395 400  
 Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly Asn Asn  
 20 405 410 415  
 Ser Glu Thr Glu Ile Phe Arg Pro Gly Gly Asp Met Arg Asp Asn  
 420 425 430  
 25 Trp Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile Glu Pro Leu  
 435 440 445  
 Gly Val Ala Pro Thr Lys Ala \*  
 30 450 455  
 (2) INFORMATION FOR SEQ ID NO:34:  
 (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 37 base pairs  
 35 (B) TYPE: Nucleic Acid  
 (C) STRANDEDNESS: Single  
 (D) TOPOLOGY: Linear  
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:34:  
 40 GGGATTCTGG ATCCAGAGCA GAAGACAGTG CCAATGA 37

(2) INFORMATION FOR SEQ ID NO:35:  
 (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 33 base pairs  
 45 (B) TYPE: Nucleic Acid  
 (C) STRANDEDNESS: Single  
 (D) TOPOLOGY: Linear  
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:35:  
 50 CTCGAGCTCC TGAAGACAGT CAGACTCATC AAG 33

(2) INFORMATION FOR SEQ ID NO:36:  
 (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 39 base pairs  
 55 (B) TYPE: Nucleic Acid  
 (C) STRANDEDNESS: Single  
 (D) TOPOLOGY: Linear  
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:36:  
 60 GGTCTAGAAG CTTTAGCCCA TAGTGCTTCC TGCTGCTCC 39

## (2) INFORMATION FOR SEQ ID NO:37:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 36 base pairs
- (B) TYPE: Nucleic Acid
- (C) STRANDEDNESS: Single
- (D) TOPOLOGY: Linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:37:

GGCGGGATCC TCGAGGTACC TGTRTGAAAGAAGCA 36

10

## (2) INFORMATION FOR SEQ ID NO:38:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 38 base pairs
- (B) TYPE: Nucleic Acid
- (C) STRANDEDNESS: Single
- (D) TOPOLOGY: Linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:38:

20

GGTCTAGAAG CTTTATGCTC CYAAGAACCC AAGGAACA 38

## (2) INFORMATION FOR SEQ ID NO:39:

25

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 7 amino acids
- (B) TYPE: Amino Acid
- (C) STRANDEDNESS: Single
- (D) TOPOLOGY: Linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:39:

Ile Gly Pro Gly Arg Ala Phe  
1 5

35

## (2) INFORMATION FOR SEQ ID NO:40:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 7 amino acids
- (B) TYPE: Amino Acid
- (C) STRANDEDNESS: Single
- (D) TOPOLOGY: Linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:40:

Ile Gly Pro Gly Arg Ala Trp  
1 5

45

## (2) INFORMATION FOR SEQ ID NO:41:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 7 amino acids
- (B) TYPE: Amino Acid
- (C) STRANDEDNESS: Single
- (D) TOPOLOGY: Linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:41:

55

Leu Gly Pro Gly Ser Thr Phe  
1 5

## (2) INFORMATION FOR SEQ ID NO:42:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 7 amino acids
- (B) TYPE: Amino Acid
- (C) STRANDEDNESS: Single
- (D) TOPOLOGY: Linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:42:

65

1 Ile Gly Pro Gly Arg Val Leu  
5

5 (2) INFORMATION FOR SEQ ID NO:43:  
(i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 7 amino acids  
(B) TYPE: Amino Acid  
(C) STRANDEDNESS: Single  
(D) TOPOLOGY: Linear  
10 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:43:

1 Ile Gly Pro Gly Ser Ala Phe  
5

15 (2) INFORMATION FOR SEQ ID NO:44:  
(i) SEQUENCE CHARACTERISTICS:  
(A) LENGTH: 5 amino acids  
(B) TYPE: Amino Acid  
(C) STRANDEDNESS: Single  
20 (D) TOPOLOGY: Linear  
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:44:

1 Ile Gly Pro Gly Arg  
5

25

## WHAT IS CLAIMED IS:

1. An isolated polypeptide comprising an HIV gp120 amino acid sequence selected from the group consisting of Sequence ID Nos. 2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, 24, 26, and 28, and fragments thereof.
2. The polypeptide of Claim 1 wherein the polypeptide additionally comprises a flag epitope sequence.
3. The polypeptide of Claim 2 wherein the flag epitope sequence is HSV gD-1 flag epitope sequence.
4. The polypeptide of Claim 2 wherein the flag epitope sequence is fused to the HIV gp120 amino acid sequence.
5. An oligonucleotide of not more than five kilobases encoding an HIV gp120 polypeptide sequence comprising an amino acid sequence selected from the group consisting of Sequence ID Nos. 2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, 24, 26, and 28, and fragments thereof.
6. The oligonucleotide of Claim 5 wherein the oligonucleotide includes a nucleotide sequence selected from the group consisting of Sequence ID Nos. 1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, and 27, and fragments thereof.
7. The oligonucleotide of Claim 5 wherein the amino acid sequence encoded by the oligonucleotide additionally comprises a flag epitope.

8. The oligonucleotide of Claim 5 wherein the flag epitope is HSV gD-1 flag epitope.
9. The oligonucleotide of Claim 7 wherein the flag epitope is fused to the HIV gp120 amino acid sequence.
10. A vaccine comprising gp120 MN and an HIV gp120 polypeptide sequence selected from the group consisting of Sequence ID Nos. 2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, 24, 26, and 28, and fragments thereof in a suitable carrier.
11. A vaccine comprising:
  - a. a first gp120 polypeptide sequence or a fragment thereof; and
  - b. a breakthrough isolate HIV gp120 polypeptide sequence or a fragment thereof from a vaccinee vaccinated with said first HIV gp120 polypeptide sequence;wherein said HIV gp120 polypeptide sequences are in a suitable carrier.
12. The vaccine of Claim 11 wherein said first HIV gp120 polypeptide sequence comprises gp120 MN, gp120 A244, gp120 MN-GNE6 (Sequence ID No. 31), or gp120 MN-GNE8 (Sequence ID No. 33).
13. The vaccine of Claim 12 wherein said vaccine additionally comprises a second gp120 polypeptide sequence comprising gp120 MN, gp120 A244, gp120 MN-GNE6 (Sequence ID No. 31), or gp120 MN-GNE8 (Sequence ID No. 33), or a fragment thereof, wherein said second HIV gp120 polypeptide sequence is different from said first HIV gp120 polypeptide sequence.

14. The vaccine of Claim 13 wherein said first gp120 polypeptide sequence comprises gp120 MN and said second gp120 polypeptide sequence comprises gp120 A244.

5

15. The vaccine of Claim 14 wherein said breakthrough isolate comprises an HIV gp120 polypeptide sequence selected from the group consisting of Sequence ID Nos. 2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, 24, 26, and 28, and fragments thereof in a suitable carrier.

16. The vaccine of Claim 13 wherein said first gp120 polypeptide sequence comprises gp120 MN and said second gp120 polypeptide sequence comprises gp120 MN-GNE8 (Sequence ID No. 33).

17. The vaccine of Claim 16 wherein said breakthrough isolate HIV gp120 polypeptide sequence is an HIV gp120 polypeptide sequence selected from the group consisting of Sequence ID Nos. 2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, 24, 26, and 28, and fragments thereof, in a suitable carrier.

25 18. The vaccine of Claim 13 wherein said breakthrough isolate HIV gp120 polypeptide is from a vaccinee vaccinated with said first and second HIV gp120 polypeptide sequences.

19. A method for making an HIV vaccine comprising adding an HIV gp120 polypeptide sequence or fragments thereof from a breakthrough isolate from a vaccinee to the vaccine with which the vaccinee was vaccinated.

5

20. The vaccine of Claim 11 wherein said first gp120 polypeptide sequence is from a macrophage-tropic HIV-1 strain.

10

21. The vaccine of Claim 11 wherein said first gp120 polypeptide sequence is from a T-cell-tropic HIV-1 strain.

15

22. The vaccine of Claim 21 wherein said vaccine additionally comprises a second gp120 polypeptide sequence or a fragment, from a macrophage-tropic HIV-1 strain.

20

23. The vaccine of Claim 22 wherein said first and second gp120 polypeptide sequences bind to different chemokine receptors.

25

24. The vaccine of Claim 23 wherein said first gp120 polypeptide sequence binds to CC-CKR-5 and said second gp120 polypeptide sequence binds to CXC-CKR-4.

30

25. The vaccine of Claim 11 wherein said vaccine additionally comprises an virus engineered to induce a cytotoxic T-cell response.

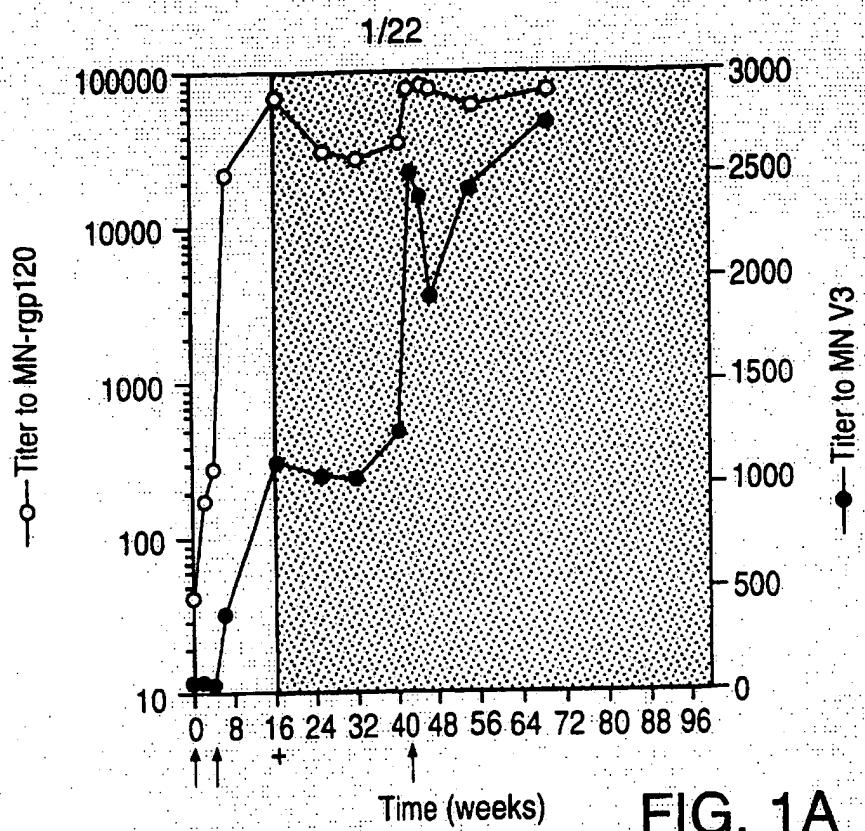


FIG. 1A

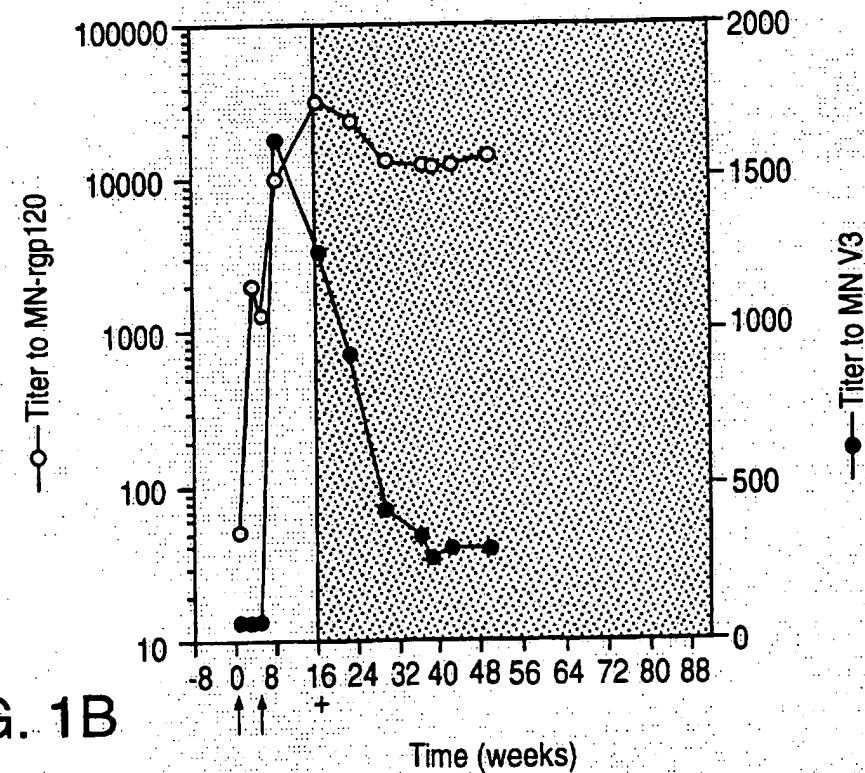


FIG. 1B

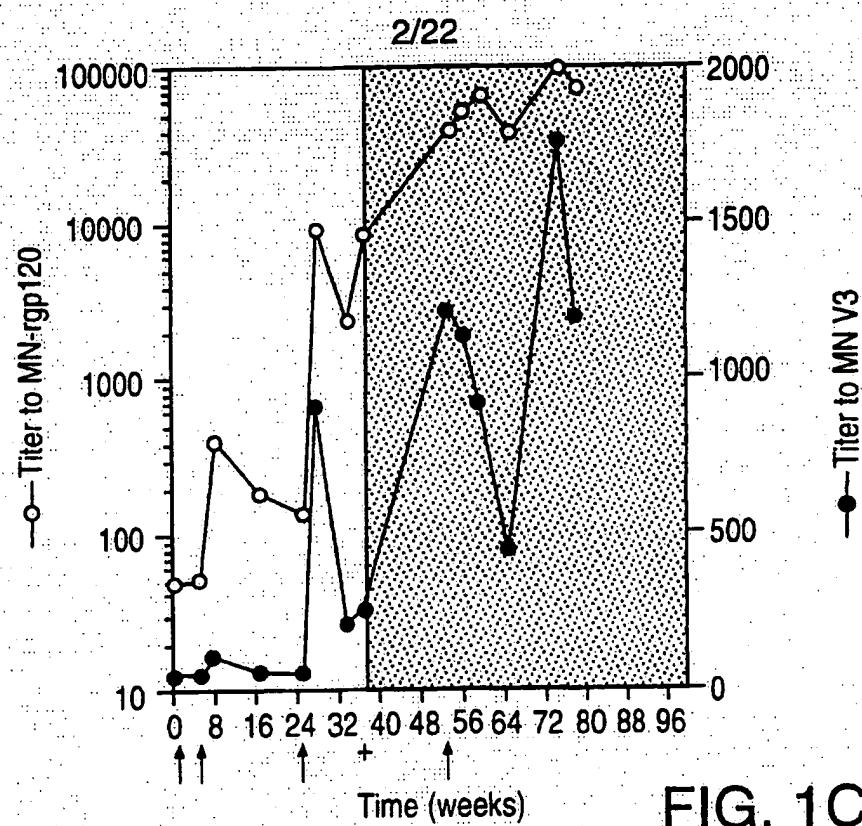


FIG. 1C

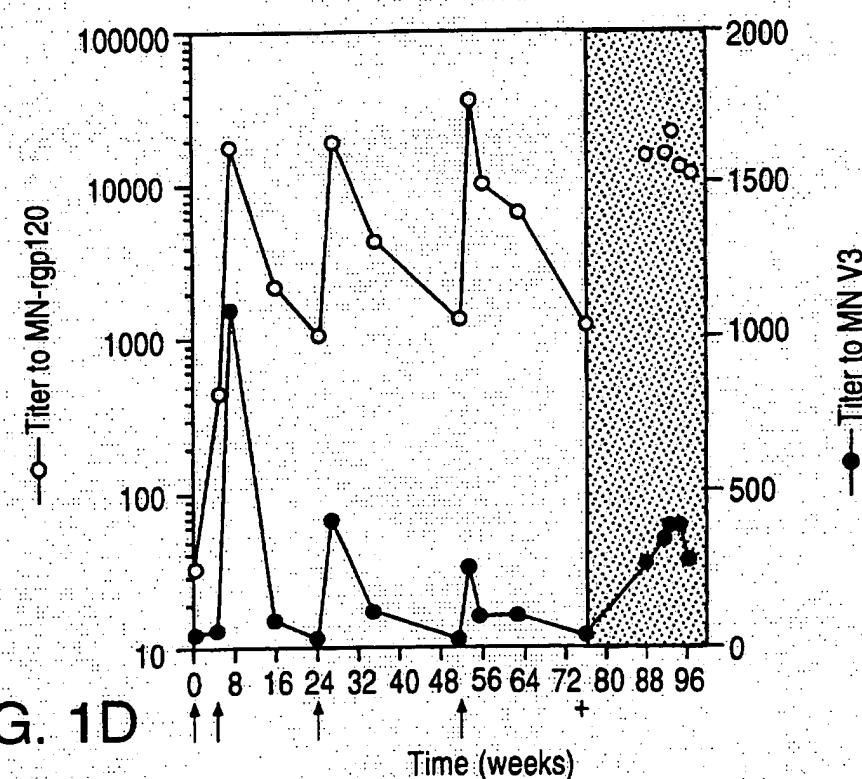


FIG. 1D

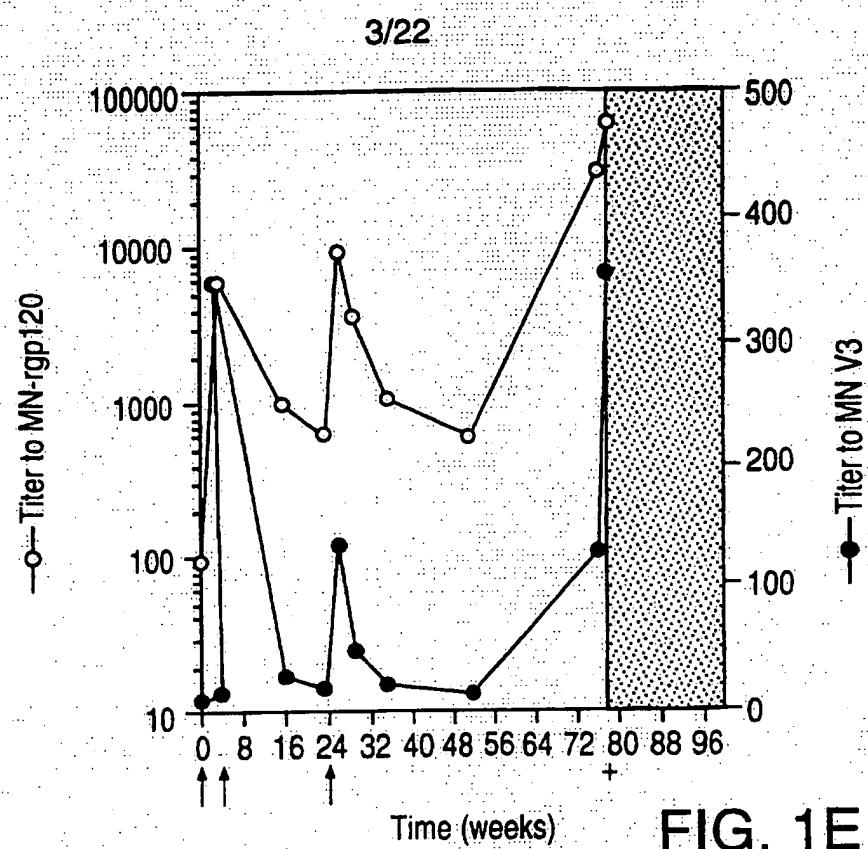


FIG. 1E

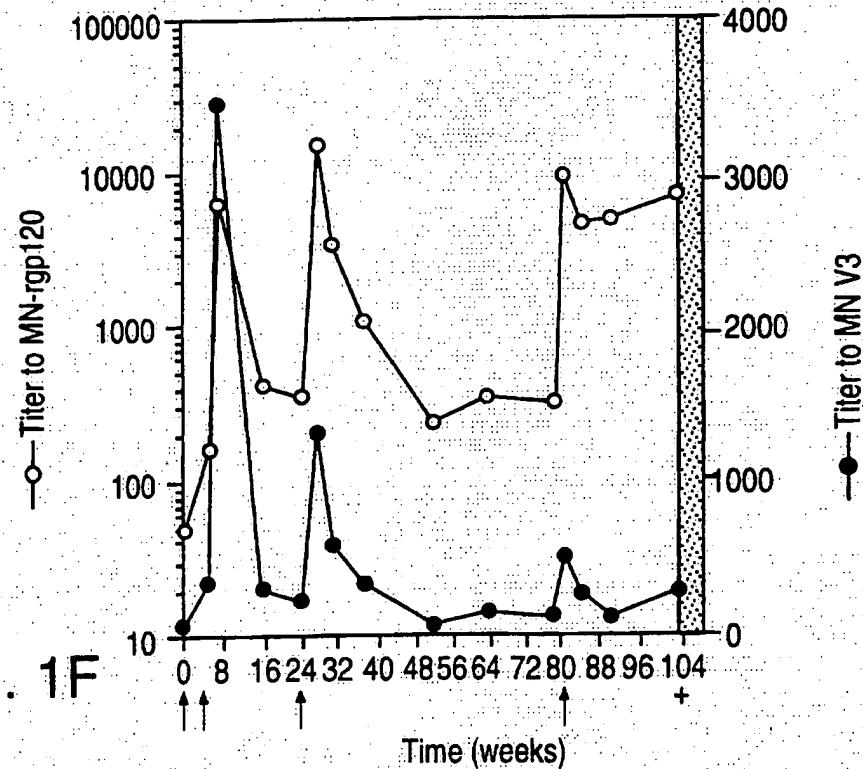


FIG. 1F

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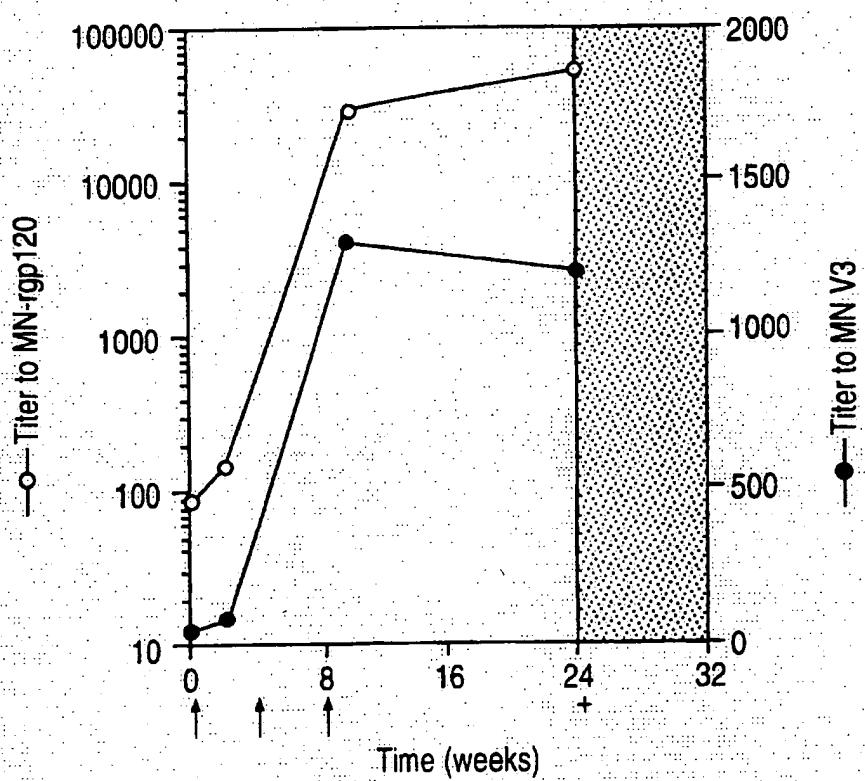


FIG. 1G

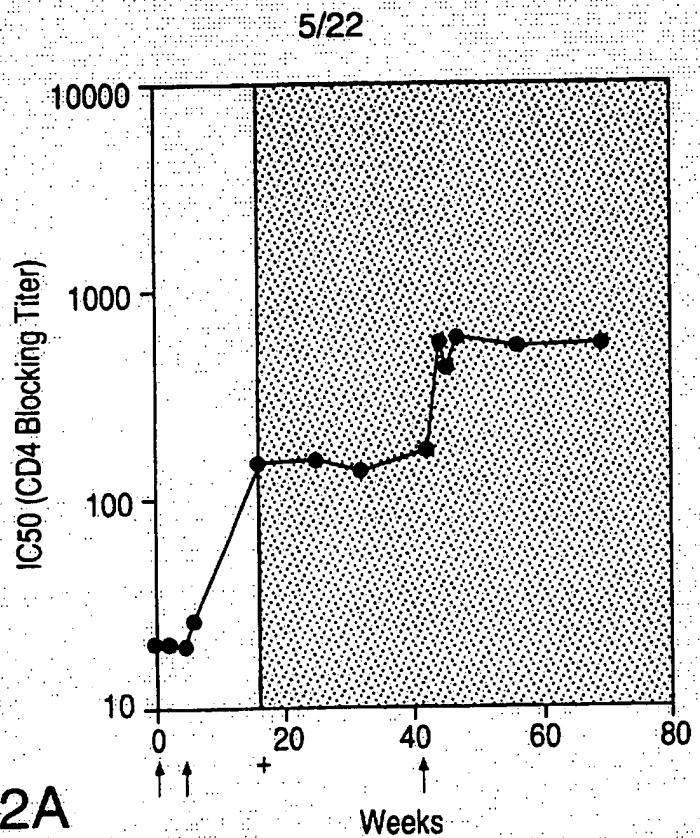


FIG. 2A

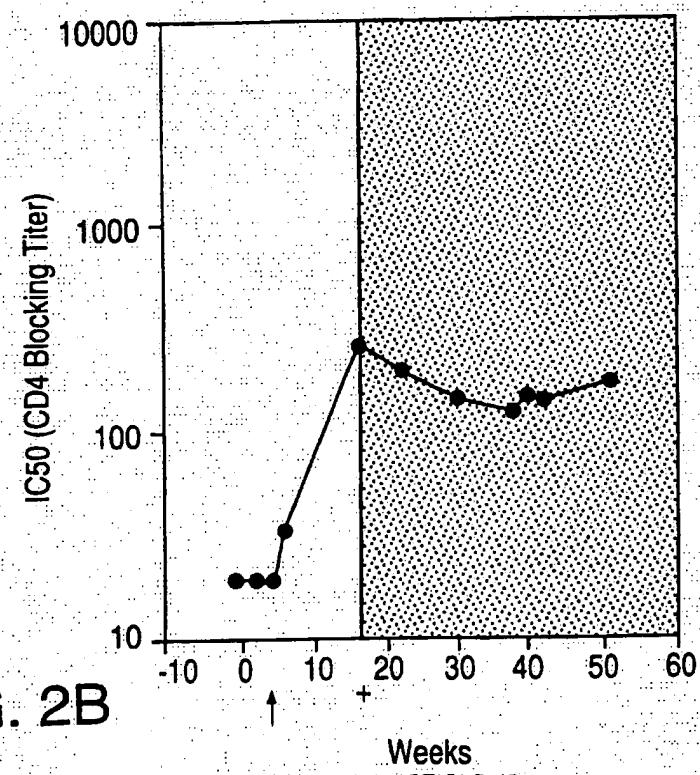


FIG. 2B

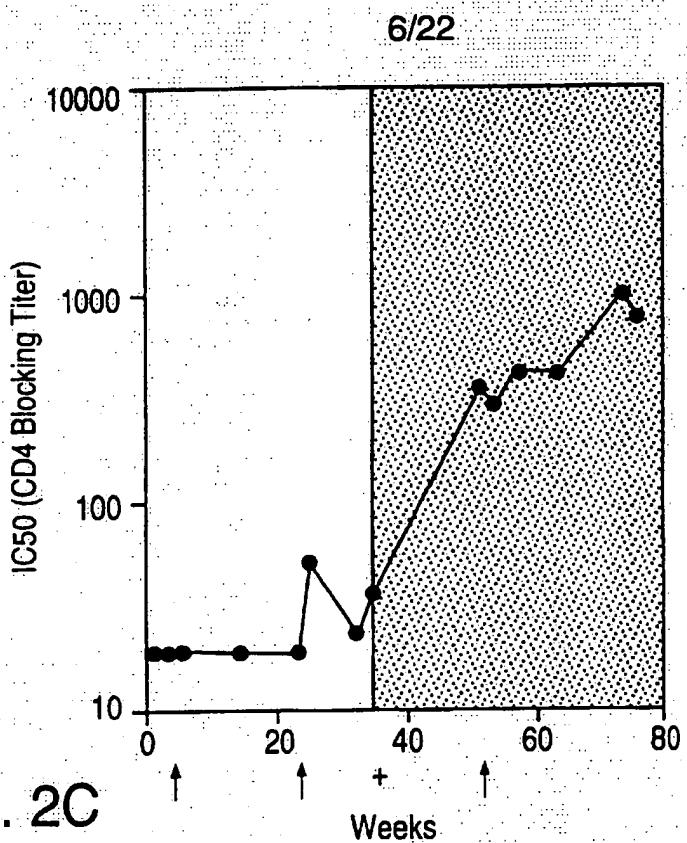


FIG. 2C

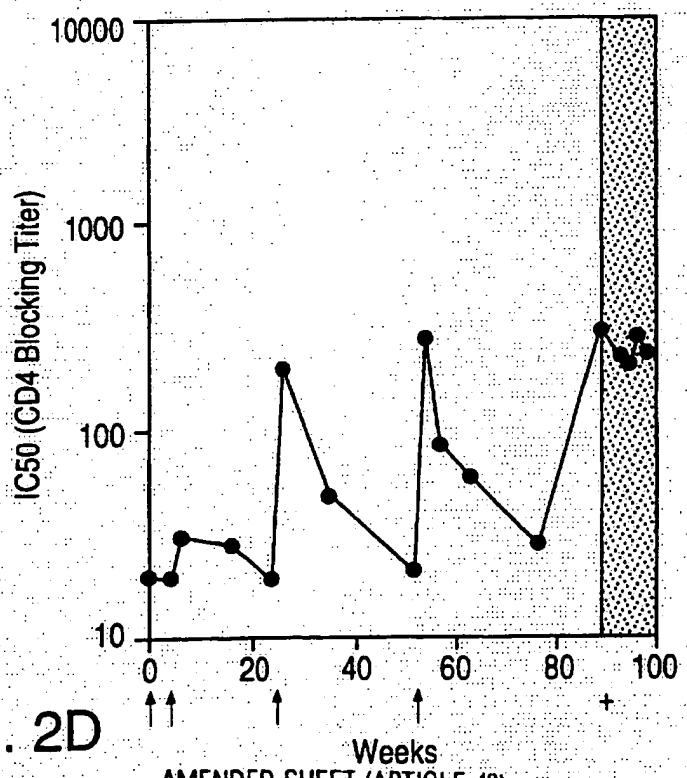


FIG. 2D

AMENDED SHEET (ARTICLE 19)

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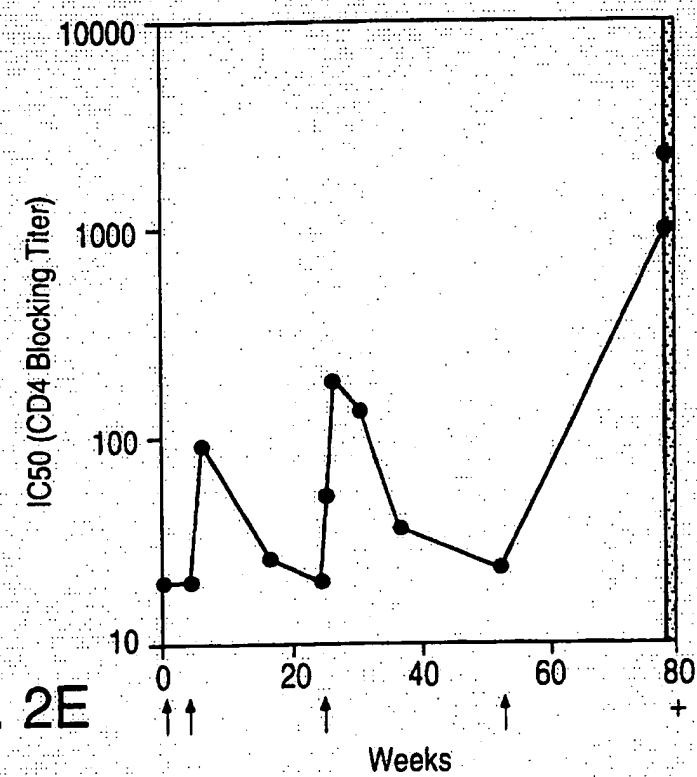


FIG. 2E

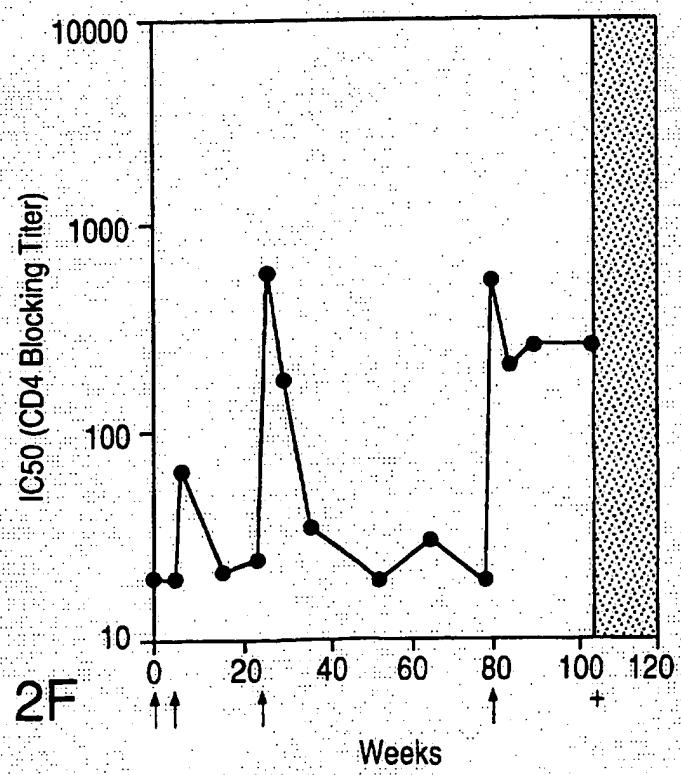


FIG. 2F

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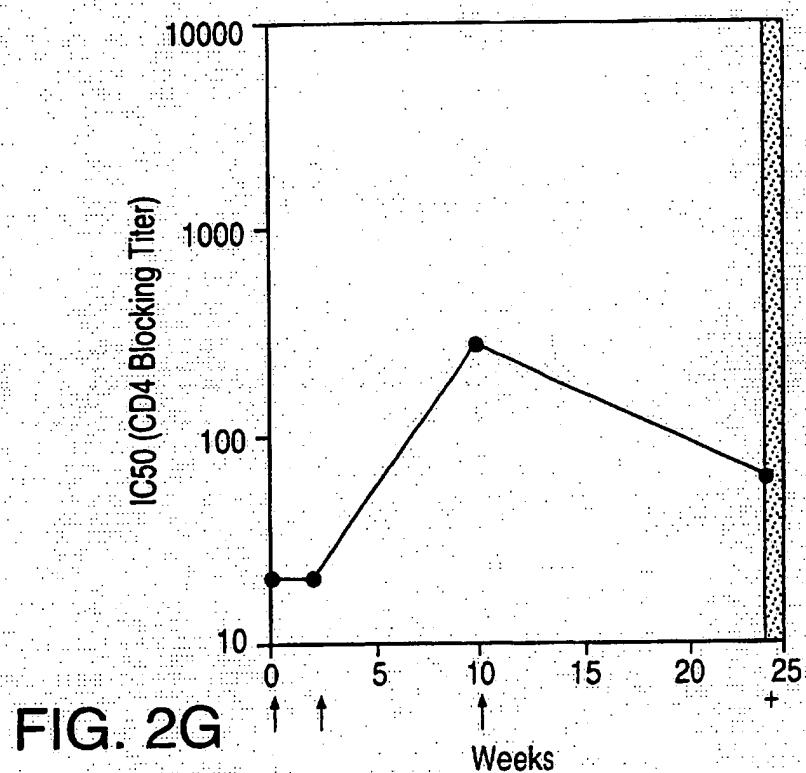


FIG. 2G

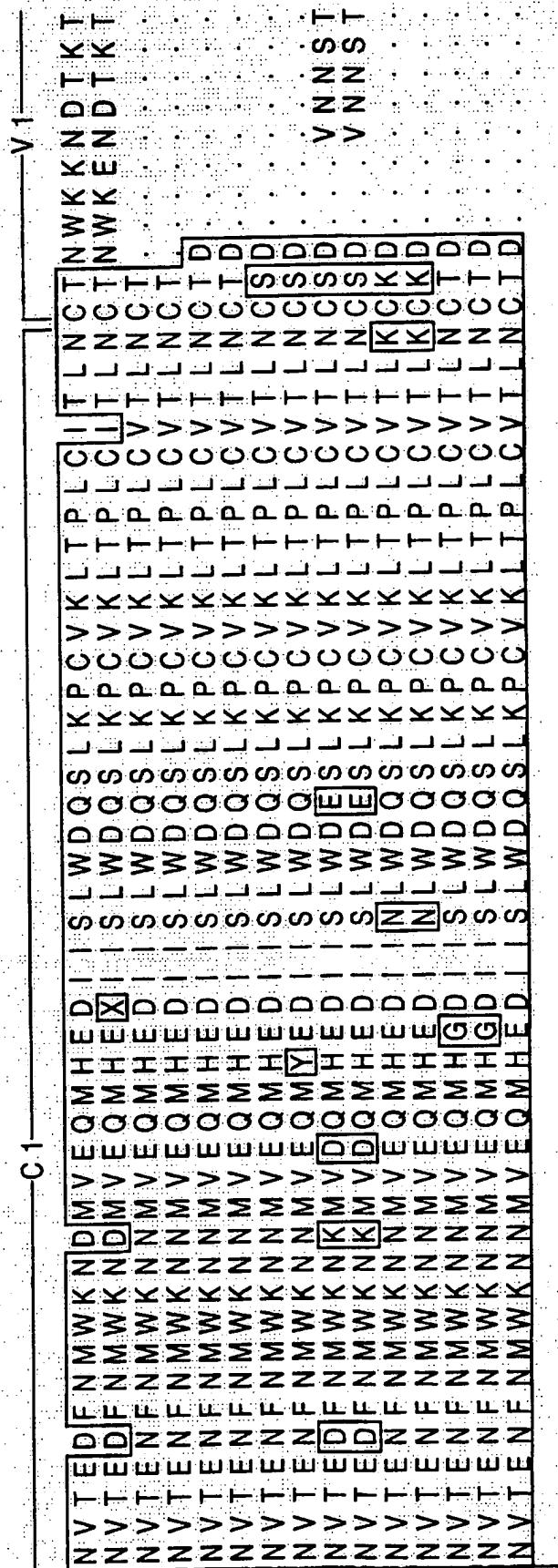
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十一

**AMENDED SHEET (ARTICLE 19)**

FIG. 3A

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AMENDED SHEET (ARTICLE 19)

FIG. 3B

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FIG. 3C

**AMENDED SHEET (ARTICLE 19)**

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FIG. 3D

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C2		C3	
240	A GFA	LK C	R DKK
240	A GFA	LK C	D KK
223	A GFA	LK C	D KK
223	A GFA	LK C	D KK
225	A GFA	LK C	D KK
225	A GFA	LK C	D KK
223	A GFA	LK C	D KK
223	A GFA	LK C	D KK
236	A GFA	LK C	D KK
236	A GFA	LK C	D KK
236	A GFA	LK C	D KK
236	A GFA	LK C	D KK
224	A GFA	LK C	D KK
224	A GFA	LK C	D KK
214	A GFA	LK C	D KK
214	A GFA	LK C	D KK
226	A GFA	LK C	D KK
C6.1			
C6.5			
C8.3			
C8.6			
C15.2			
C15.3			
C7.2			
C7.10			
C115			
C11.7			
C105			
C107			
C17.1			
C17.3			
MNGNE			

FIG. 3E

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37

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**AMENDED SHEET (ARTICLE 19)**

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FIG. 3G

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FIG. 3

FIG. 31

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三

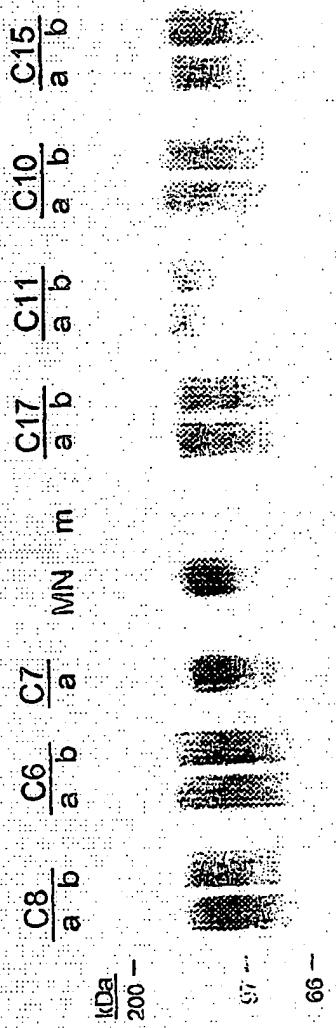
**AMENDED SHEET (ARTICLE 19)**

FIG. 3J

FIG. 3A	FIG. 3B
FIG. 3C	FIG. 3D
FIG. 3E	FIG. 3F
FIG. 3G	FIG. 3H
FIG. 3I	FIG. 3J

33  
E

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AMENDED SHEET (ARTICLE 19)

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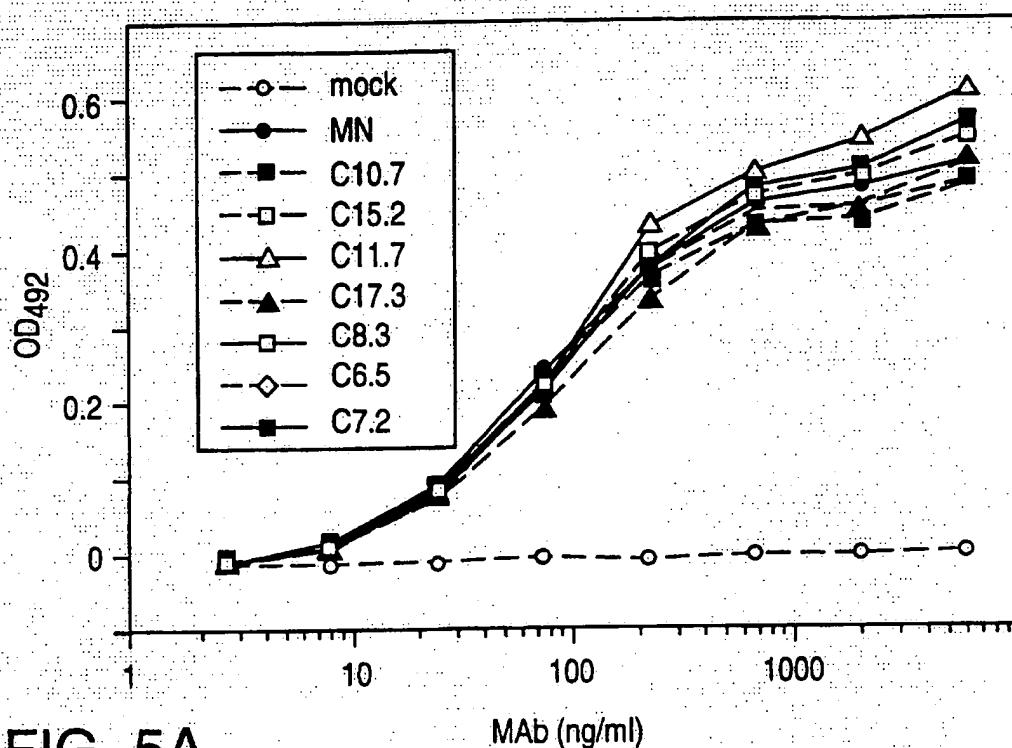


FIG. 5A

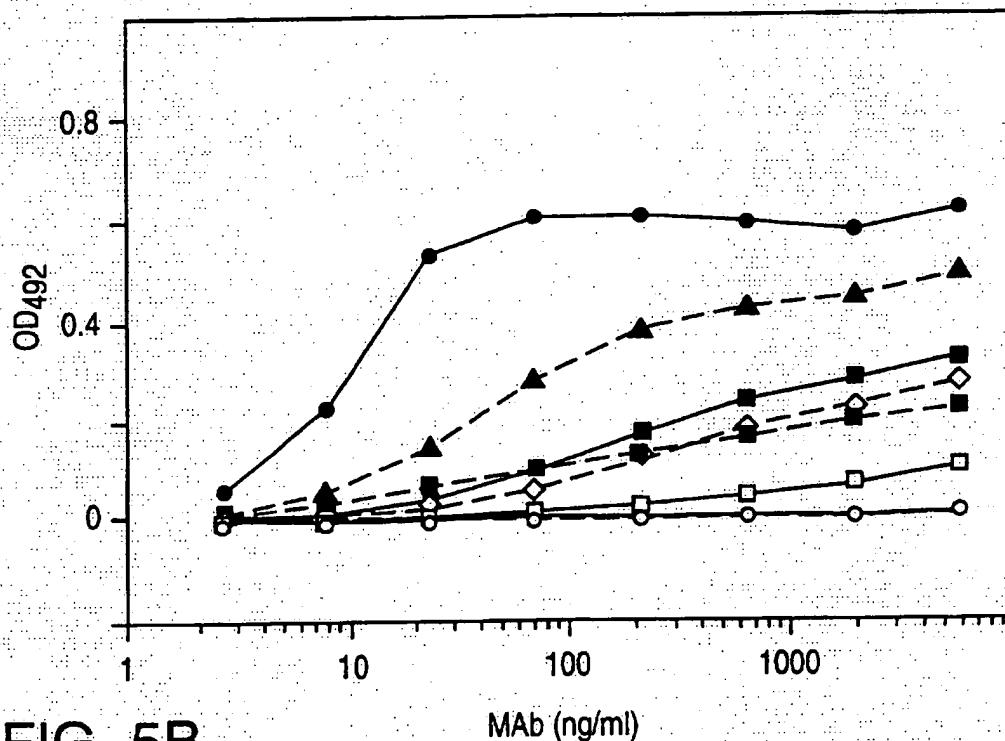


FIG. 5B

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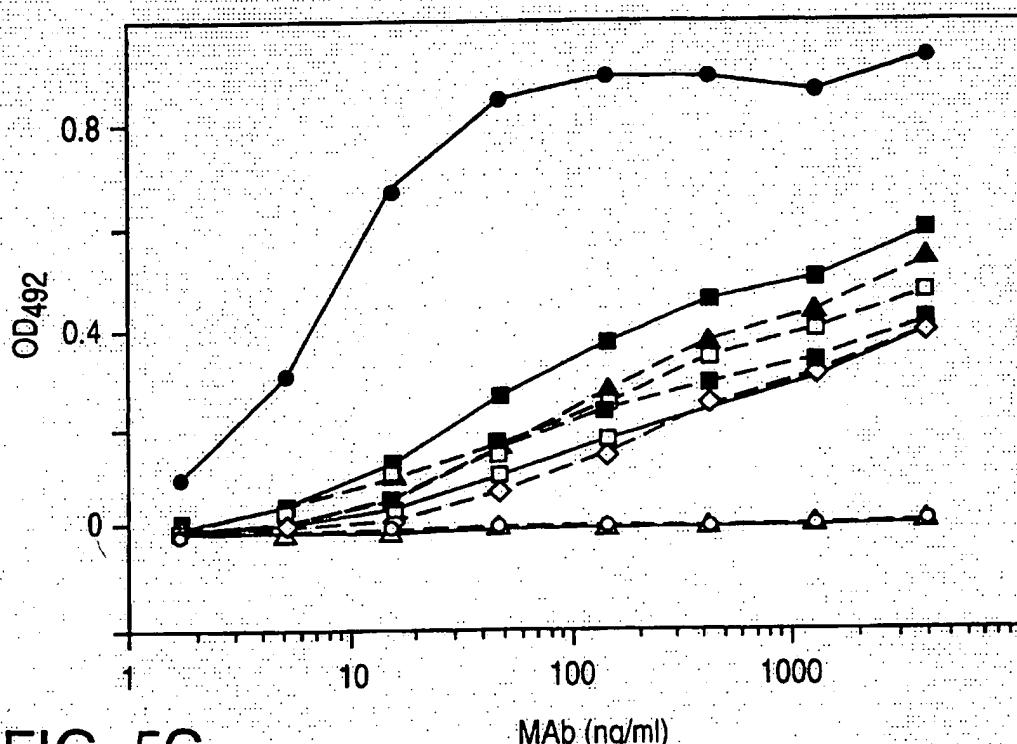


FIG. 5C

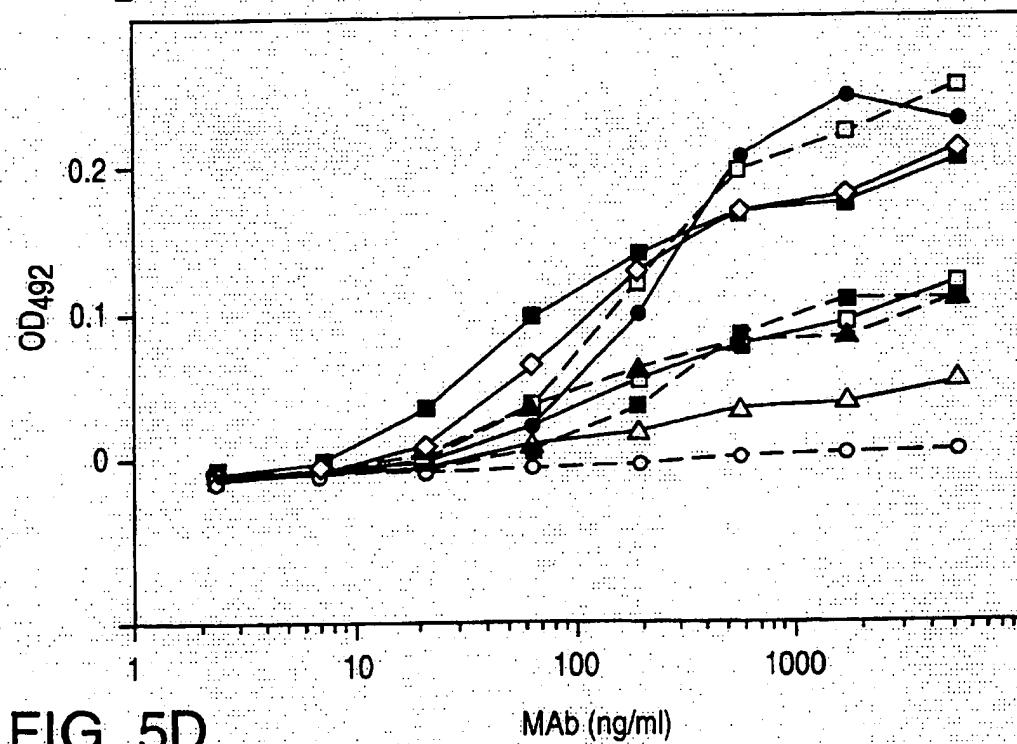


FIG. 5D

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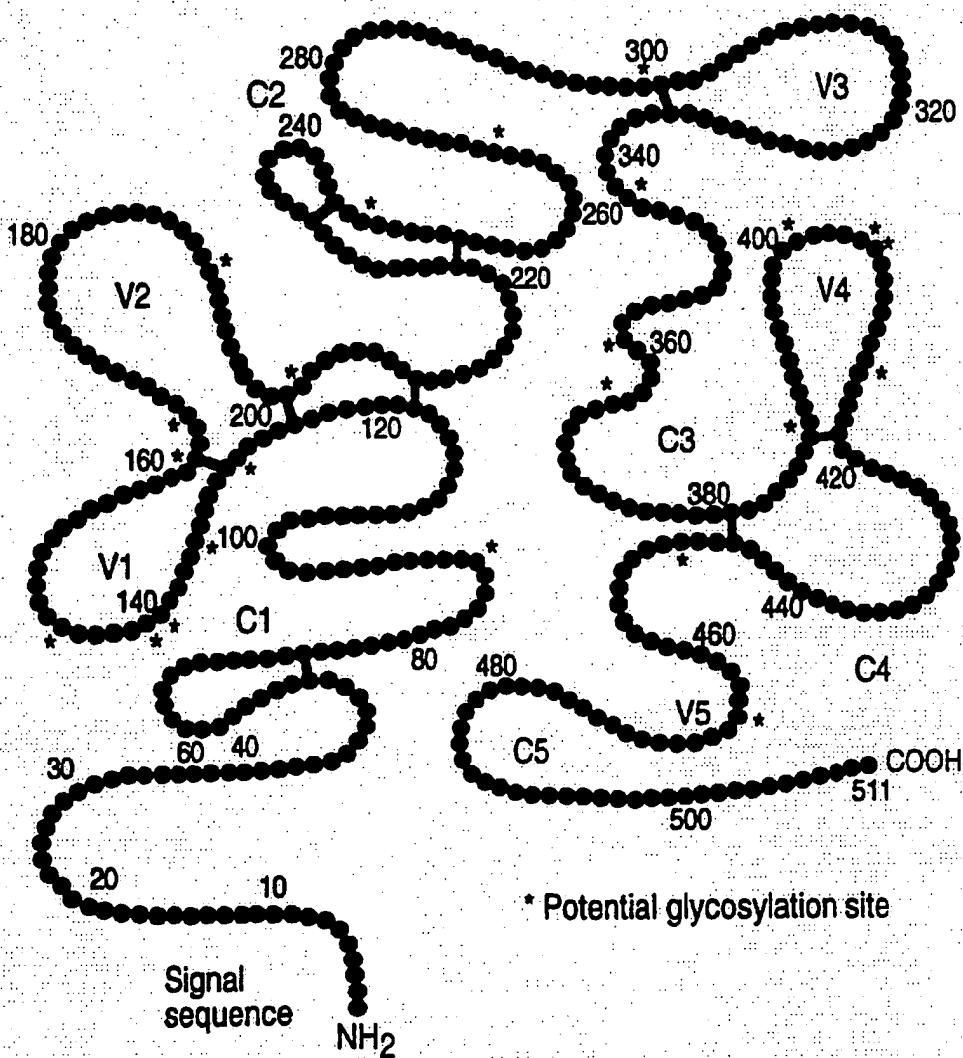


FIG. 6

# INTERNATIONAL SEARCH REPORT

International Application No.

PCT/US 97/09690

**A. CLASSIFICATION OF SUBJECT MATTER**  
**IPC 6 C12N15/49 C07K14/16 A61K39/21**

According to International Patent Classification (IPC) or to both national classification and IPC

**B. FIELDS SEARCHED**

Minimum documentation searched (classification system followed by classification symbols)

**IPC 6 C12N C07K A61K**

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 94 28929 A (GENETECH, INC.) 22 December 1994 see page 56 SEQ. ID. NO. 25. see page 50, line 14. - line 31 ---	1-18, 20-25
X	P.W. BERMAN ET AL.: "Genetic and immunologic characterization of viruses infecting MN-rgp120 vaccinated volunteers" ONE WORLD, ONE HOPE: XI INTERNATIONAL CONFERENCE ON AIDS, vol. 10, no. supplement 3, 7 - 12 July 1996, VANCOUVER, CANADA, page 10 XP002045307 See "Methods" in Abstract Mo.A.285 ---	1,5,6 -/-

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

\* Special categories of cited documents:

- \*A\* document defining the general state of the art which is not considered to be of particular relevance
- \*E\* earlier document but published on or after the international filing date
- \*L\* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- \*O\* document referring to an oral disclosure, use, exhibition or other means
- \*P\* document published prior to the international filing date but later than the priority date claimed

\*T\* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

\*X\* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

\*Y\* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

\*&\* document member of the same patent family

Date of the actual completion of the international search

30 October 1997

Date of mailing of the international search report

26-11-1997

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2  
NL - 2280 HV Rijswijk  
Tel: (+31-70) 340-2040, Tx: 31 651 apo nl  
Fax: (+31-70) 340-3016

Authorized officer

Cupido, M

# INTERNATIONAL SEARCH REPORT

International Application No.

PCT/US 97/09690

**C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT**

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	<p>M.J. MCELRATH ET AL.: "Human immunodeficiency virus type 1 infection despite prior immunization with a recombinant envelope vaccine regimen" PROCEEDINGS OF THE NATIONAL ACADEMY OF SCIENCES OF USA, vol. 93, no. 9, 30 April 1996, WASHINGTON US, pages 3972-3977, XP002045308 see page 3976, last paragraph; figure 1</p> <p>---</p>	19
T	<p>P.W. BERMAN ET AL.: "Genetic and immunologic characterization of viruses infecting MN-rgp120-vaccinated volunteers" THE JOURNAL OF INFECTIOUS DISEASES, vol. 176, no. 2, August 1997, pages 384-397, XP002045309 see the whole document</p> <p>-----</p>	1-25

# INTERNATIONAL SEARCH REPORT

## Information on patent family members

International Application No.

PCT/US 97/09690

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